

# REPORT DOCUMENTATION PAGE

*Form Approved  
OMB No. 074-0188*

Public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing this collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to Washington Headquarters Services, Directorate for Information Operations and Reports, 1215 Jefferson Davis Highway, Suite 1204, Arlington, VA 22202-4302, and to the Office of Management and Budget, Paperwork Reduction Project (0704-0188), Washington, DC 20503.

<b>1. AGENCY USE ONLY (Leave blank)</b>			<b>2. REPORT DATE</b> Sept 96	<b>3. REPORT TYPE AND DATES COVERED</b> Technical Report.	
<b>4. TITLE AND SUBTITLE</b>  Laboratory Scale Stabilization of N-Springs Groundwater Strontium-90 Using Phosphatic Materials			<b>5. FUNDING NUMBERS</b>  N/A		
<b>6. AUTHOR(S)</b> T.E. Moody, S.W. Petersen, E.G. Torne, J. Vicakova, & J.F. Higginbotham					
<b>7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES)</b>  CH2M HILL Hanford, Inc.  Oregon State University Radiation Center			<b>8. PERFORMING ORGANIZATION REPORT NUMBER</b> BHI-00864 Rev.0  <b>19980710 103</b>		
<b>9. SPONSORING / MONITORING AGENCY NAME(S) AND ADDRESS(ES)</b>  SERDP 901 North Stuart St. Suite 303 Arlington, VA 22203			<b>10. SPONSORING / MONITORING AGENCY REPORT NUMBER</b>  N/A		
<b>11. SUPPLEMENTARY NOTES</b> Report prepared for the US DOE Office of Environmental Restoration and Waste Management. This work was supported in part by DOE.					
<b>12a. DISTRIBUTION / AVAILABILITY STATEMENT</b>  Approved for public release: distribution is unlimited					<b>12b. DISTRIBUTION CODE</b> A
<b>13. ABSTRACT (Maximum 200 Words)</b>  This study was initiated to investigate the potential use of phosphatic materials as permeable geochemical barriers for groundwater contaminated with strontium-90. Groundwater discharges to the Columbia River create potential human food chain hazards. It is imperative to immobilize the contamination before it reaches the river. Phosphate materials have been proven by various researchers to be chemical compounds that combine with contaminant metals forming into insoluble metal-phosphate minerals. These minerals are stable and insoluble under normal soil conditions. The US DOE is currently undertaking phosphate stabilization projects at Hanford, Oak Ridge, Savannah, Fernald, and the University of Idaho.					
<b>14. SUBJECT TERMS</b> Strontium-90, Groundwater contamination, Phosphates, SERDP					<b>15. NUMBER OF PAGES</b> 111
					<b>16. PRICE CODE</b> N/A
<b>17. SECURITY CLASSIFICATION OF REPORT</b> unclass.	<b>18. SECURITY CLASSIFICATION OF THIS PAGE</b> unclass.	<b>19. SECURITY CLASSIFICATION OF ABSTRACT</b> unclass.	<b>20. LIMITATION OF ABSTRACT</b> UL		

NSN 7540-01-280-5500

Standard Form 298 (Rev. 2-89)  
Prescribed by ANSI Std. Z39-18  
298-102

DTIC QUALITY INSPECTED 1

BHI-00864  
Rev. 0

# **Laboratory Scale Stabilization of N-Springs Groundwater Strontium-90 Using Phosphatic Materials**



Prepared for the U.S. Department of Energy  
Office of Environmental Restoration and  
Waste Management

**Bechtel Hanford, Inc.**  
Richland, Washington

**TRADEMARK DISCLAIMER**

Reference herein to any specific commercial product, process, or service by trade name, trademark, manufacturer, or otherwise, does not necessarily constitute or imply its endorsement, recommendation, or favoring by the United States Government or any agency thereof or its contractors or subcontractors.

This report has been reproduced from the best available copy.  
Available in paper copy and microfiche.

Available to the U.S. Department of Energy  
and its contractors from  
Office of Scientific and Technical Information  
P.O. Box 62  
Oak Ridge, TN 37831  
(615) 576-8401

Available to the public from the U.S. Department of Commerce  
National Technical Information Service  
5285 Port Royal Road  
Springfield, VA 22161  
(703) 487-4650

Printed in the United States of America

DISCLM-5.CHP (8-91)

BHI-00864  
REV: 0  
OU: N/A  
TSD: N/A  
ERA: N/A

### APPROVAL PAGE

Title of Document: Laboratory Scale Stabilization of N-Springs Groundwater Strontium-90  
Using Phosphatic Materials

Authors: T. E. Moody                    E. G. Torne  
             S. W. Petersen                    J. Vlcakova  
    J. F. Higginbotham

Approval: J. D. Isaacs, Groundwater Project Engineer

R. P. Schroeder for J. D. ISAACS  
Signature

9-26-96  
Date

G. C. Henckel, Manager  
Groundwater Project

  
Signature

9/26/96  
Date

The approval signature on this page indicates that this document has been authorized for issue to U.S. Department of Energy, Richland Operations Office through appropriate channels. No other forms or signatures are required to document this information release.

BHI-DIS 9-27-96 MH

BHI-00864  
Rev. 0

# Laboratory Scale Stabilization of N-Springs Groundwater Strontium-90 Using Phosphatic Materials

**Authors**

T. E. Moody

S. W. Petersen

CH2M HILL Hanford, Inc.

E. G. Torne

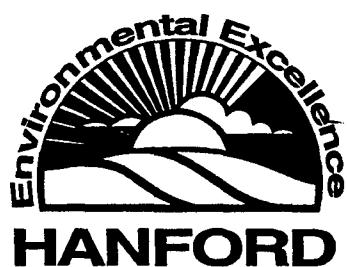
J. Vlcakova

J. F. Higginbotham

Oregon State University Radiation Center

**Date Published**

September 1996



Prepared for the U.S. Department of Energy  
Office of Environmental Restoration and  
Waste Management

**Bechtel Hanford, Inc.**

Richland, Washington

## EXECUTIVE SUMMARY

This study was initiated to investigate the potential use of phosphatic materials as permeable geochemical barriers for groundwater contaminated with strontium-90 ( $^{90}\text{Sr}$ ). Groundwater discharges to the Columbia River create potential human food chain hazards. It is imperative to immobilize the contamination before it reaches the river. Phosphate materials have been proven by various researchers to be chemical compounds that combine with contaminant metals forming into insoluble metal-phosphate minerals. These minerals are stable and insoluble under normal soil conditions. The U.S. Department of Energy is currently undertaking phosphate stabilization projects at Hanford, Oak Ridge, Savannah, Fernald, and the University of Idaho.

The objective of this study was to demonstrate the precipitation, efficiency of formation, amounts of phosphatic material required for precipitation, and the stability of a strontium-phosphate mineral phase. This bench-scale engineering study will serve to establish the chemical needed for a nominal barrier configuration. The phosphate material reacts with  $^{90}\text{Sr}$  in the passing groundwater to create a stable and insoluble strontiapatite mineral.

The soil used in this bench-scale experiment was taken from a depth of 51 to 53 ft in a  $^{90}\text{Sr}$ -contaminated well (N-Springs well 199-N-105A). The soil contains 500 pCi/g  $^{90}\text{Sr}$  along with smaller amounts of  $^{137}\text{Cs}$ ,  $^{152}\text{Eu}$ ,  $^{60}\text{Co}$ , and  $^3\text{H}$ . Heavy metal contamination includes arsenic, cadmium, lead, and selenium above background concentrations.

Batch desorption experiments were conducted to determine the maximum  $^{90}\text{Sr}$  concentration as a function of time. Batch adsorption isotherm experiments were conducted on the soil by adding incremental amounts of phosphate material below and above the amounts needed for strontiapatite formation as a function of mineral phase molar ratios. Column tests using  $^{90}\text{Sr}$  contaminated soil were performed to examine the pH stability of the formed strontiapatite mineral.

The results indicate that bone char and hydroxyapatite are efficient at sorbing  $^{90}\text{Sr}$  from soil and groundwater. The North Carolina apatite failed to be an efficient sorbent material due to its high strontium content. A fourth material, Ash Meadows clinoptilolite (hereafter referred to as clinoptilolite), was also evaluated and proved to be effective in sorbing  $^{90}\text{Sr}$  desorbed from soil.

Results from this study indicate that phosphate materials can be effective as permeable barriers for reducing  $^{90}\text{Sr}$  concentrations from soil and groundwater.

## CONTENTS

1.0	INTRODUCTION .....	1-1
2.0	APATITE AND PHOSPHATIC MATERIALS .....	2-1
3.0	DESCRIPTION OF EXPERIMENTS .....	3-1
3.1	DESORPTION AND ADSORPTION EXPERIMENT .....	3-1
3.2	KINETICS OF COMPLEXATION .....	3-2
3.3	LOADING CAPACITY OF SORBENT MATERIALS .....	3-3
3.4	pH STABILITY TEST OF PRECIPITATED <sup>90</sup> Sr-PHOSPHATE MINERALS .....	3-3
3.5	ANALYTICAL METHODS .....	3-4
3.5.1	Liquid Scintillation Counting .....	3-4
3.5.2	Atomic Absorption Spectrophotometry .....	3-4
3.5.3	ICP-MS Analysis .....	3-4
3.5.4	Ion Chromatography .....	3-4
4.0	RESULTS AND DISCUSSION .....	4-1
4.1	ASSUMPTIONS OF THE METHODOLOGY .....	4-1
4.2	DESORPTION ISOTHERM .....	4-1
4.3	ADSORPTION ISOTHERMS .....	4-2
4.4	KINETICS OF COMPLEXATION .....	4-5
4.5	LOADING CAPACITY EXPERIMENT .....	4-6
4.6	pH STABILITY TEST OF <sup>90</sup> Sr-PHOSPHATE COMPLEX .....	4-7
4.7	SUMMARY OF RESULTS .....	4-8
4.7.1	Adsorption Isotherms .....	4-8
4.7.2	Kinetics of Complexation .....	4-8
4.7.3	Loading Capacity Experiment .....	4-9
4.7.4	pH Stability Test of <sup>90</sup> Sr-Phosphate Complex .....	4-9
5.0	REFERENCES .....	5-1

## FIGURES

4-1.	Desorption of Strontium-90 from N-Springs Soil, Well 199-N-105A, Using Hanford Simulated Groundwater .....	4-11
4-2.	Reduction of Solution <sup>90</sup> Sr with the Addition of Incremental Amounts of Sorbent During 72 Hours Contact Time .....	4-12
4-3.	Plot of Normalized Abundances of Major Element Compositions in Contact with Various Sorbent Materials .....	4-13

4-4.	Changes in the Anion Composition of the Liquid Phase Illustrated by Normalized Element Concentrations .....	4-14
4-5.	Plot of the Weight Fraction of HAP Versus the Mass Excess (Adsorbed Solute) for Alkali and Earth Alkali Elements .....	4-15
4-6.	Time Required for Strontium-90 Sorption Using a 10:1 Water-to-Soil Ratio and a Fixed Amount of Sorbent Material .....	4-16
4-7.	Strontium-90 Loading Capacity of the Sorbents Expressed as $K_d$ Values .....	4-17
4-8.	Strontium-90 in Solution After Complexed with Sorbent for 1, 2, and 4 Weeks .....	4-18
4-9.	pH Stability of the Incubated 1, 2, and 4 Week $^{90}\text{Sr}$ -Sorbent Complexes .....	4-19

## TABLES

3-1.	List of the Adsorbent Materials and Related Properties .....	3-6
3-2.	Concentration of the Solutes in Simulated Hanford Groundwater .....	3-6
3-3.	Concentration of Elements in the Standard Solutions Used for Atomic Absorption Spectrophotometry .....	3-7
3-4.	Concentration of Elements in the Standard Solutions Used for Ion Chromatography .....	3-7

## APPENDICES

A.	Experiment A: Primary Adsorption Experiment .....	A-i
B.	Experiment B: Kinetics of Complexation .....	B-i
C.	Experiment C: Loading Capacity of Sorbents .....	C-i
D.	Experiment D: pH Stability Test .....	D-i

**ACRONYMS**

AAS	atomic absorption spectroscopy
clinoptilolite	Ash Meadows clinoptilolite
EMPS	electron microprobe spectrometry
HAP	hydroxyapatite
IC	ion chromatography
ICP	inductively coupled plasma-mass spectroscopy
INAA	instrumental neutron activation analysis
LSC	liquid scintillation counting
MS	mass spectroscopy
NC apatite	North Carolina apatite
REE	rare earth elements

BHI-00864  
Rev. 0

## 1.0 INTRODUCTION

This report presents the results of a laboratory study designed to evaluate the ability of phosphatic materials to sorb strontium-90 ( $^{90}\text{Sr}$ ) from soil and groundwater. This investigation was initiated in response to an initiative to evaluate new technologies for use in remedial activities at the Hanford Site. The phosphatic materials were evaluated relative to Ash Meadows clinoptilolite (hereafter referred to as clinoptilolite), a current ion exchange medium used in the remediation of  $^{90}\text{Sr}$  contaminated groundwater.

The radionuclide  $^{90}\text{Sr}$  is produced by spontaneous fission of uranium-233 ( $^{233}\text{U}$ ) and plutonium-239 ( $^{239}\text{Pu}$ ), which were produced in the reactors at Hanford. The yield of  $^{90}\text{Sr}$  from these radionuclides is 6.9 percent and 2.11 percent, respectively. Elevated levels of  $^{90}\text{Sr}$  in soil and groundwater are a concern because the element strontium behaves in a manner very similar to calcium in chemical and biological systems. Calcium is an important element in most fauna and is easily adsorbed by organisms from water and other ingested materials. Therefore, it is desirable to control the release of radioactive strontium before it enters the biological and biogeochemical cycle where it can be incorporated in bone and dental tissue. Conversely, the same mechanism that allows  $^{90}\text{Sr}$  to incorporate into bone will be examined to test the efficacy of phosphatic materials to stabilize  $^{90}\text{Sr}$ .

Previous studies (e.g., Lee, 1961; Anderson, 1971; Moody and Wright, 1995; Carter, 1990) have shown that phosphate materials are effective in precipitating several heavy metals including Pb, Pu, Sr, and U. Phosphate compounds have the ability to incorporate these and other elements into their crystal structure forming a stable, insoluble compound resistant to remobilization by inorganic or organic processes. The half life of  $^{90}\text{Sr}$  is 29.1 years. If  $^{90}\text{Sr}$  could be effectively complexed and immobilized for several hundred years, elevated concentrations would naturally decay to insignificant levels. The incorporation of  $^{90}\text{Sr}$  into a calcium phosphate mineral will possibly result in the formation of strontiapatite, a mineral that is insoluble under normal environmental conditions.

This study used soil from the well 199-N-105A in the 100-N Area. The soil was recovered from a depth of 51 to 53 ft (15.5 to 16 m), known to have a very high concentration of  $^{90}\text{Sr}$ . The source of  $^{90}\text{Sr}$  was the 1301-N and 1325-N cribs. The concentration of  $^{90}\text{Sr}$  at this soil depth was approximately 500 pCi/g. This is the highest  $^{90}\text{Sr}$  concentration for 100-N Area borehole sediments (Serne and LeGore, 1996). The soil was used to test the capacity of various phosphate compounds to precipitate  $^{90}\text{Sr}$  and investigate the subsequent stability of the phosphate materials under various physical conditions.

BHI-00864  
Rev. 0

## 2.0 APATITE AND PHOSPHATIC MATERIALS

Phosphatic minerals are important accessory minerals in igneous, metamorphic, and sedimentary rocks. Although typically constituting less than 1 percent of common rocks, phosphate minerals can be a major rock forming phase. For example, sedimentary phosphate rock is dominated by 90 to 95 percent apatite.

The most abundant mineral of the phosphate group is apatite with the chemical composition  $\text{Ca}_5(\text{PO}_4)_3(\text{OH},\text{F},\text{Cl})$ . There is a complete isomorphous solid solution between the end-members hydroxyapatite,  $\text{Ca}_5(\text{PO}_4)_3(\text{OH})$ , fluorapatite,  $\text{Ca}_5(\text{PO}_4)_3(\text{F})$ , and chlorapatite  $\text{Ca}_5(\text{PO}_4)_3(\text{Cl})$ . Apatite is an important mineral for environmental remediation because it has a low solubility and a propensity for incorporation of heavy metals into its crystal structure. A recent review of apatite solubility by Chandler and Fuerstenau (1984) indicated a  $\text{pK}_s$  value of 115. The solubility of apatite with strontium in place of calcium (strontiapatite) is reported to be  $\log K_{sp} = -51.3$ . (Verbeeck, et al., 1977). In field application, strontium movement in soils has been stabilized by the use of phosphate amendments (Anderson, 1971; Francis, 1978). Because of its low tendency to dissolve, it may be one of the most suitable minerals considered for retardation of radionuclides from groundwater.

Various trace elements are readily incorporated into the apatite crystal structure (Liu and Comodi, 1993). Hughes et al. (1991) evaluated the substitution of rare earth elements (REE) into apatite with emphasis on the structural crystallography. Elements of the alkali-earth group are also very compatible with some apatite minerals. For example, fluorapatite can contain over 11 molecular percent SrO (Deer et al., 1992). Although this is a rather unusually high concentration, Edgar (1989) pointed out that many analyses of apatite do not include SrO or BaO, so the actual concentrations of these elements are unknown in many instances.

The crystal structure of apatite belongs to the hexagonal centrosymmetric space group  $P6_3/m$ . The structure of the lattice can also be degraded to monoclinic symmetries in some chlorapatites when certain anion solid solutions are present (Hughes et al., 1989, 1991). There are two sites for metallic cations: Ca1 which represents a calcium site with six neighboring oxygen atoms (six fold coordination) and Ca2, a calcium site with eight oxygen and one F, Cl, or OH anion (nine fold coordination). Isomorphous substitution can occur in both Ca1 and Ca2 sites by Na, K, Sr, Mg, Ba, Zn, Cd, Sc, Y, the REEs and U.

BHI-00864  
Rev. 0

### 3.0 DESCRIPTION OF EXPERIMENTS

The experiments were designed to test various geochemical and kinetic responses of different types of sorbent materials in contact with  $^{90}\text{Sr}$  in solution. These sorbent materials are listed in Table 3-1. Although not presented in the table, phosphorus pentoxide ( $\text{P}_2\text{O}_5$ ) and monocalcium phosphate ( $\text{Ca}(\text{H}_2\text{PO}_4)_2 \cdot \text{H}_2\text{O}$ ) were tested. The phosphate materials chosen were found by previous studies to be effective in scavenging metals from soil and groundwater. In addition to the natural phosphates, such as North Carolina apatite (NC apatite) and bone char, pure, reagent-grade hydroxyapatite (HAP) was also studied under the same conditions as the phosphate materials to evaluate effects caused by impurities in the natural apatite. A series of experiments were also done with clinoptilolite. This mineral was evaluated to compare the chemical effectiveness of phosphatic material as a geochemical barrier to a current methodology in groundwater contaminant retardation.

After receiving the  $^{90}\text{Sr}$  contaminated soil from well 199-N-105A, the soil was air dried, mixed thoroughly, and sieved using a 2 mm ASTM sieve. This was done to ensure a uniform  $^{90}\text{Sr}$  concentration in the subsequent experiments.

The experiments as outlined in the scope of work consisted of four discreet steps:

1. The desorption/adsorption study, which focused on the geochemical characterization of simulated groundwater solutions that were in contact with the soil and sorbent material.
2. The kinetics study, which quantified the rate of the sorption reactions.
3. The loading capacity study, which determined the amount of  $^{90}\text{Sr}$  adsorbed by different phosphatic materials.
4. The pH stability test for the precipitated  $^{90}\text{Sr}$ -phosphate complexes, which examined the stability of different pH conditions and time periods.

#### 3.1 DESORPTION AND ADSORPTION EXPERIMENT

In this portion of the bench scale study, a measured amount of Hanford soil and Hanford simulated groundwater were mixed with incremental amounts of different phosphate materials. When groundwater is in contact with solid mineral phases in soils, its composition is dependent on the initial groundwater concentration, adsorbed and precipitated solutes, desorbed constituents, and dissolved crystals. In order to characterize the groundwater after equilibration, aliquots of the solution were analyzed after varying agitation periods with the soil phosphate mixtures.

A series of preliminary experiments were performed to identify the appropriate amounts of soil and phosphatic materials needed for determining of isotherms. The optimal times for

equilibration of the soil and groundwater were also determined in this phase of the study. Results of these tests are presented in Tables A-3 and A-4. Results indicate that 3 g of soil reacted for 72 hours were sufficient for determining adsorption isotherms and dissolution/precipitation reactions. It was also determined that proportions of sorbent > 10 percent by weight generally did not increase adsorption. Therefore, the range of interest was limited to 0 to 10 percent by weight of adsorbent, and various intervals between these bounds were used for the primary study.

Each of these experiments used  $3.00 \pm 0.01$  g of Hanford soil and 30 mL of Hanford simulated groundwater. The groundwater solution was prepared from double distilled water by adding reagent grade chemicals (Table 3-2) to reproduce the major element abundance of natural groundwater in Hanford soils. In this way changes in the major element composition of groundwater can be related to the sorbent added.

Incremental amounts of sorbent materials were added to the soil and groundwater mixture to determine adsorption isotherms and dissolution/precipitation reactions. A centrifuge tube of each series was agitated for 72 hours in an automatic shaker. The soil was separated from the groundwater by centrifugation. A syringe equipped with a nucleopore filter membrane was used to withdraw the solution. The conductivity and pH were determined at this point using a conductivity meter and pH electrode. Aliquots of the solution were analyzed for  $^{90}\text{Sr}$  by liquid scintillation counting (LSC) spectroscopy, for major cationic constituents by atomic absorption spectroscopy (AAS), for minor elements by ICP-MS, and for anion abundances by ion chromatography (IC).

Statistical and analytical uncertainties that occur in any type of batch experiment were avoided by running each series in duplicate. The assumption that the composition of soil and sorbent in each of the vessels is homogeneous was made considering the uniform grain size achieved by sieving the samples. Another assumption is that the reaction containers, which consisted of 60 mL polyethylene tubes, were inert to the system. However,  $^{90}\text{Sr}$  and other trace elements could also have been adsorbed or precipitated on surfaces of containers. Because the addition of a strong dissociating mineral acid could not be accomplished in the agitation tube, it is assumed that sorption of the wall may be uniform in all experiments. To minimize these effects in the storage containers that were used to transport groundwater to the analytical instruments, the solution was acidified with dilute ICP-MS grade nitric acid.

### 3.2 KINETICS OF COMPLEXATION

Adsorption reactions are typically rapid and equilibrate in the order of minutes to hours (Sposito, 1989; Moody and Wright, 1995). Therefore the time intervals for this experiment were chosen to be 1, 3, 5, 10, 15, and 24 hours. Considering the most effective concentration of the sorbent, the composition of the soil/sorbent mixture was selected from the primary adsorption experiment.

Each centrifuge tube was filled with 3 g of soil, the appropriate amount of sorbent material, and 30 mL Hanford simulated groundwater. The vessels were then shaken for the desired agitation

time, the solid phase was separated by centrifugation for 30 minutes, and the supernatant was filtered through a 0.45  $\mu\text{m}$  nucleopore membrane filter.

Conductivity and pH were measured immediately after filtration on 3 mL of the solution. Aliquots of the solution were then analyzed for  $^{90}\text{Sr}$  by LSC, major cation components by AAS, and trace element cations by ICP-MS.

### **3.3 LOADING CAPACITY OF SORBENT MATERIALS**

This experiment was performed to quantify the maximum amount of strontium in solution that can be adsorbed by the different sorbent materials. Stock solutions containing concentrations of 0.01, 0.1, 1, and 3 g of strontium nitrate per 30 mL double distilled water were prepared by weighing the appropriate amounts in polyethylene bottles and adding double distilled water.

Centrifuge tubes were filled with 3 g of NC apatite, bone char, hydroxyapatite and clinoptilolite. To each of these tubes 30 mL of the stock solution was added. The solutions were agitated for two days and then filtered through a 0.45  $\mu\text{m}$  nucleopore membrane filter. The concentration of strontium was then analyzed by AAS.

### **3.4 pH STABILITY TEST OF PRECIPITATED $^{90}\text{Sr}$ -PHOSPHATE MINERALS**

The stability of the  $^{90}\text{Sr}$ -phosphate was investigated under various pH conditions and reaction times as outlined in the scope of work. The soil was mixed with phosphate and clinoptilolite at proportions that were most effective for the stabilization of the contaminant. A total of 1 kg soil-sorbent mix was used for each experimental condition.

Three replicates of each mixture were prepared by saturating them with Hanford simulated groundwater and allowing the sample to dry to field capacity (0.33 bar). Each of the samples were then incubated for 3 different time periods: 1, 2, and 4 weeks. After incubation, an aliquot of the soil-sorbent mixture was removed and placed in a column. Six columns were prepared to study the column water flow-through for each treatment at pH levels of 5, 6, 7, 8, and 9. A control column that contained soil without added sorbent material was prepared for each of the treatments. A volume of Hanford simulated groundwater equal to 5 pore volumes of material and adjusted for the proper pH was introduced into the columns, then reacted water was analyzed.

During the last phase of this experiment, a saturation extract was performed on 200 g of soil from each column. The soil was saturated with groundwater and the liquid phase was extracted and analyzed for  $^{90}\text{Sr}$ , ICP-MS metals, IC anions, pH and conductivity.

### 3.5 ANALYTICAL METHODS

All of the analyses were performed on the liquid phase. The concentration of  $^{90}\text{Sr}$  was measured by LSC, major cations in the groundwater by AAS, minor constituents by ICP-MS, anions by IC and pH and conductivity by electrode probe.

#### 3.5.1 Liquid Scintillation Counting

A volume of 2 mL per sample was mixed with 15 mL of commercial scintillation cocktail (OPTIFLOUR) in 20 mL liquid scintillation glass vials. Prior to counting,  $^{90}\text{Sr}$  was equilibrated with its daughter product  $^{90}\text{Y}$  for two weeks. A Beckman<sup>TM</sup> LS 6500 liquid scintillation counter was used for the measurement. Blank samples containing 2 mL of distilled water and cocktail were used to correct for background.

#### 3.5.2 Atomic Absorption Spectrophotometry

Sodium, magnesium, and calcium concentrations in the liquid phase were determined by the AAS method. A Perkin Elmer Model 5000 spectrophotometer was used. The relatively high abundance of these elements in the solution made a dilution to lower concentrations necessary. Dilution of the samples was carried out by adding 0.2 mL of 70 percent  $\text{HNO}_3$ , 0.41 mL of Cs (1000 ppm), and 14.8 mL of double distilled water to exactly 5 mL of the original sample. The dilution process had to be repeated more than 4 times for some of the samples.

The concentration of elements in the standard solutions is listed in Table 3-3. The standards were run at the beginning, middle, and end of each analytical series. The number of unknowns that were analyzed in sequence did not exceed 22.

#### 3.5.3 ICP-MS Analysis

Five milliliter aliquots of the groundwater solutions were acidified with 0.7 percent  $\text{HNO}_3$  to prevent precipitation of solutes on the container walls. An internal standard containing Be, In, and Bi was added to the sample. Standards were prepared from ICP-MS grade solutions. A standard was analyzed for quality control for every eleventh sample of unknown composition. A VG Plasma Quad ICP-MS<sup>TM</sup> system manufactured by Fisons was used.

#### 3.5.4 Ion Chromatography

The analysis of anions in the groundwater was performed by ion chromatography on an DIONEX<sup>TM</sup> 2000i ion chromatograph located in the Oregon State University College of Agricultural Sciences. Aliquots of 2-5 mL of the unknown solution were used. The columns consisted of AG4A-SC guard and AS4A separator material. The regeneration of the columns was accomplished by elution with 1.8 molar  $\text{Na}_2\text{CO}_3$  and 1.7 molar  $\text{NaHCO}_3$  solutions. The solutions were run through the columns at 2 mL/minute.

Standards were prepared from DIONEX™ anion standard stock solutions (Lot # 950623). They contained fluorine, chlorine, nitrate, phosphate, and sulfate. Five standard solutions were prepared (Table 3-4). The accuracy of the analysis was ascertained by repeated runs of standard solution 3. An analysis of that standard was performed after every twelfth unknown sample.

**Table 3-1. List of the Adsorbent Materials and Related Properties**

<b>Property</b>	<b>NC Apatite</b>	<b>Bone Char</b>	<b>Hydroxyapatite</b>	<b>Clinoptilolite</b>
Formula	$\text{Ca}_{10}(\text{PO}_4)_6(\text{OH},\text{F})_2$	$\text{Ca}_{10}(\text{PO}_4)_6(\text{OH},\text{F})_2$	$\text{Ca}_{10}(\text{PO}_4)_6(\text{OH},\text{F})_2$	zeolite
Origin	phosphate rock	charred animal bone	synthetic chemical	aggregate
Grain Size	180-250 $\mu\text{m}$	180-250 $\mu\text{m}$	180-250 $\mu\text{m}$	180-250 $\mu\text{m}$
Surface	smooth	porous	not available	porous
Texture	rounded	irregular fragments	plates	rounded
Analysis	INAA and EMPS	INAA and EMPS	INAA	INAA

The surface morphology was examined by scanning electron microscopy and petrographic thin section observation. Chemical analysis were performed by instrument neutron activation analysis (INAA) and electron microprobe spectroscopy (EMPS).

**Table 3-2. Concentration of the Solutes in Simulated Hanford Groundwater**

<b>Element</b>	<b>Conc. (mmol/L)</b>	<b>Weight (mg/L)</b>	<b><math>\text{CaCO}_3</math> (mmol/L)</b>	<b><math>\text{MgCO}_3</math> (mmol/L)</b>	<b><math>\text{NaHCO}_3</math> (mmol/L)</b>	<b><math>\text{CaSO}_4</math> (mmol/L)</b>	<b><math>\text{Ca}(\text{NO}_3)_2</math> (mmol/L)</b>
$\text{Ca}^{2+}$	0.36	14.5	0.12	-	-	0.16	0.08
$\text{Mg}^{2+}$	0.18	4.5	-	0.19	-	-	-
$\text{Na}^+$	0.13	3	-	-	0.13	-	-
$\text{SO}_4^{2-}$	0.16	15.7	-	-	-	0.16	-
$\text{NO}_3^-$	0.16	10	-	-	-	-	0.16
$\text{HCO}_3^-$	0.87	53.1	0.37	0.38	0.13	-	-

The pH of this solution was  $8.2 \pm 0.1$ .

**Table 3-3. Concentration of Elements in the Standard Solutions Used for Atomic Absorption Spectrophotometry**

Element	Standard 1 $\mu\text{g/mL}$	Standard 2 $\mu\text{g/mL}$	Standard 3 $\mu\text{g/mL}$	Standard 4 $\mu\text{g/mL}$	Standard 5 $\mu\text{g/mL}$	Standard 6 $\mu\text{g/mL}$
Sodium	2	4	8	16	24	NA
Calcium	0.32	0.96	1.92	2.88	3.84	5
Magnesium	0.16	0.48	0.96	1.44	1.92	NA

NA = Not applicable

**Table 3-4. Concentration of Elements in the Standard Solutions Used for Ion Chromatography**

Element	Standard 1 $\mu\text{g/mL}$	Standard 2 $\mu\text{g/mL}$	Standard 3 $\mu\text{g/mL}$	Standard 4 $\mu\text{g/mL}$	Standard 5 $\mu\text{g/mL}$
Fluoride	0.20	0.51	1.02	1.53	2.04
Chloride	0.30	0.75	1.51	2.26	3.01
Nitrate	1.01	2.53	5.05	7.58	10.10
Phosphate	1.52	3.80	7.60	11.40	15.20
Sulfate	1.52	3.80	7.60	11.40	15.20

BHI-00864

Rev. 0

## 4.0 RESULTS AND DISCUSSION

Summaries of the results of the four experiments are presented in this section. The data from all of the experiments are contained in Appendices A through D.

### 4.1 ASSUMPTIONS OF THE METHODOLOGY

After receiving the  $^{90}\text{Sr}$  contaminated soil from well 199-N-105A, the soil was air dried, mixed thoroughly, and sieved using a 2 mm ASTM sieve. This was done to ensure a uniform  $^{90}\text{Sr}$  concentration in the subsequent experiments. The assumption was made that each of the centrifuge tubes contained a representative sample. Any variations in the composition of the soils would result in errors in the measurements. It should be noted that this assumption may not be valid. The precision of the analytical data was higher than the data points reproduced from duplicate runs. Because only two duplicate analysis were performed in most of the runs, it is statistically unwarranted to calculate the standard deviation of the paired observations. At least 4 to 5 repeated analysis may have to be carried out in order to confidently calculate the errors. A modified experimental approach would be a better solution to the problem of the geological heterogeneity of the samples. However, this approach was not covered in the contractor scope of work. A method to accomplish this is currently under development.

### 4.2 DESORPTION ISOTHERM

In an effort to determine the amount of  $^{90}\text{Sr}$  desorbed from the solid phase during a specific time, a desorption isotherm was executed on the Hanford soil. Two separate desorption isotherms were performed during a seven-day period. Two water-to-soil ratios were used in an effort to determine the effect of varying ratios: the standard water-to-soil ratio of 10:1 and a 5:1 water-to-soil ratio. The results of these two desorption isotherms are presented in Figure 4-1. The analytical data are presented in Table A-1. As seen from the desorption isotherm, the maximum amount of  $^{90}\text{Sr}$  desorbed from the soil into the Hanford simulated groundwater is approximately 11,000 pCi/L for the 10:1 water-to-soil ratio isotherm, and occurs at the 72 hour desorption time. For the 5:1 water-to-soil ratio isotherm, the maximum desorbed amount of 12,900 pCi/L was also 72 hours. This is only a 17 percent increase in desorbed  $^{90}\text{Sr}$  relative to twice the amount of soil in contact with solution. Data from Table A-1 can be used to calculate the kinetic rate constant  $K_a$ , assuming a first order reaction, expressed by the relationship:

$$\log\left(1 - \frac{q_t}{q_e}\right) = -k_a \times t \quad (1)$$

where

$q_t$  is the amount of  $^{90}\text{Sr}$  adsorbed on the soil

$q_e$  the concentration at time t  
 $k_d$  is the desorption rate coefficient

The experimental results of data from Table A-1 were used to calculate linear regressions. For the experiment with 3 g soil per 30 mL groundwater the desorption rate constant is 0.04/day; whereas, the system representing 6 g of soils is only 0.03/day. This is contrary to the expectation that twice as much sediment will yield double the activity.

An important question is whether  $^{90}\text{Sr}$  is actually quantitatively desorbed from the minerals or if the process is only partially completed. The  $^{90}\text{Sr}$  concentration in the soil must be known in order to evaluate the hypothesis. Because the activity of the radionuclide is 500 pCi/g, the amount of  $^{90}\text{Sr}$  per gram of soil can be calculated as follows:

$$N = \frac{(-dN/dt)}{\lambda} = \frac{-18.5[\text{Bq/g}]}{7.58 \times 10^9 [\text{sec}^{-1}]} = 2.44 \text{nuclei/g} \quad (2)$$

This concentration can also be expressed in number of isotopes per gram and would be 0.04 pmol/g. Hence a contaminated area measuring 1 x 100 x 100 m contains approximately 9.7 grams of  $^{90}\text{Sr}$ . Three grams of soil contain 55.5 Becquerel radiation. However, only in the desorption experiments, between 8.1 and 12 Becquerel of radiation was found in 30 mL of simulated groundwater after one week. Therefore, it can be concluded that only about 21 percent of  $^{90}\text{Sr}$  will partition in the liquid phase.

Considering the results of the desorption isotherm, 72 hours will be used as the amount of contact time for the adsorption isotherms. The water-to-soil ratio of 10:1 will be used for the execution of the adsorption isotherm.

#### 4.3 ADSORPTION ISOTHERMS

Adsorption isotherms are important tools for investigating precipitation and adsorption effects of solutes added to a specific solution in increasing amounts. The adsorption isotherm graphically illustrates the precipitation or reduction of the soluble metals when the sorbent is applied to the system. Adsorption is most often described in terms of isotherms, which show the relationship between the effective concentration in solution of the species being adsorbed and the actual amount adsorbed at a constant temperature. When plotted, the shape and the mathematical expression of the isotherm provides a great deal of information concerning the chemistry and sorption mechanisms within the system. The most effective concentration of the sorbent to effect precipitation can be ascertained by examining the sharp downward curves of the isotherm. The system can be modeled thermodynamically by first analyzing the solution components, entering the data into the computer program, and changing the amounts of sorbent in the theoretical system to match those amounts from the adsorption isotherm. The thermodynamic modeling results will be presented in a forthcoming document.

Two adsorption isotherms were determined. A preliminary adsorption isotherm was carried out to find the approximate range of sorbents required to induce precipitation and/or adsorption. The data for the preliminary adsorption isotherm are presented in Tables A-3 and A-4.

Hydroxyapatite, bone char, and clinoptilolite effectively reduce the solution concentration of  $^{90}\text{Sr}$ . The addition of the phosphate compounds  $\text{Ca}(\text{H}_2\text{PO}_4)_2 \cdot \text{H}_2\text{O}$  and  $\text{P}_2\text{O}_5$  in the preliminary adsorption isotherm were found to liberate the soil  $^{90}\text{Sr}$  into solution. This was undoubtedly due to the pH lowering effect of these two compounds. As a consequence, these two phosphatic compounds were excluded from further experimental consideration. Data collected from the preliminary adsorption isotherm indicates that maximum reduction of  $^{90}\text{Sr}$  in solution occurs at sorbent concentrations of less than 10 percent (mass basis). The primary adsorption isotherm was executed using 10 percent as the upper limit of the sorbent concentration. To illustrate distinct departure from linearity and to effectively determine the concentration of sorbent giving rise to the precipitation/adsorption of  $^{90}\text{Sr}$  from solution, the number of increasing concentrations of sorbent was increased from the 4 concentrations used in the preliminary adsorption isotherm. Table A-5 presents the concentrations of the added sorbent and the resulting concentration of  $^{90}\text{Sr}$  remaining in solution after 72 hours of contact. The adsorption isotherms for NC apatite, hydroxyapatite, bone char, and clinoptilolite are presented in Figure 4-2. The preliminary adsorption isotherm 20 percent sorbent value for  $^{90}\text{Sr}$  is presented in Figure 4-2 to add completeness to the linearity.

The starting  $^{90}\text{Sr}$  concentration for all of the sorbents (0 percent) is fixed at 12,111 pCi/L (determined from the desorption isotherm. Based on the 72 hour adsorption isotherm, the clinoptilolite appears to have the greatest ability to reduce  $^{90}\text{Sr}$  solution concentration. The greatest reduction occurs at clinoptilolite values of less than 2 percent. The amount of  $^{90}\text{Sr}$  precipitated and/or adsorbed at 2 percent added clinoptilolite is approximately 85 percent. The bone char is second in efficiency in reducing the  $^{90}\text{Sr}$  concentration. All concentrations of bone char reduce  $^{90}\text{Sr}$  concentration. However, 85 percent reduction in  $^{90}\text{Sr}$  concentration is achieved at higher concentrations of bone char: between 10 and 20 percent. At 20 percent added bone char the  $^{90}\text{Sr}$  concentration is nearly the same as the reduction of  $^{90}\text{Sr}$  caused by the addition of clinoptilolite. This data shows that bone char can effectively reduce the concentration of  $^{90}\text{Sr}$  (as does the clinoptilolite) but at higher values of added bone char.

The remainder of the data generated for the adsorption isotherm experiments are contained in Tables A-5 thru A-7. Data for the major cations calcium, magnesium, and sodium were recalculated to molar abundances, normalized to 100 percent, and plotted in the ternary diagram in Figure 4-3. Hanford groundwater, shown by the hatched circle in the figure, has subequal parts of sodium and calcium with 10 to 15 molar percent magnesium. When NC apatite and hydroxyapatite were added to the system, compositions became enriched in calcium relative to sodium and magnesium. Therefore it can be concluded that small amounts of calcium were introduced by adding apatite. It could be speculated, that to some degree, calcium on the surfaces was exchanged for other cations in the groundwater, but this should be supported by further microscopic study.

Bone char and clinoptilolite behaved differently with respect to sodium and magnesium concentrations. The clinoptilolite added only sodium to the groundwater; whereas, bone char

contributed sodium and calcium. The argument can be made for clinoptilolite, that ion exchange may be the prevailing mechanism to desorb sodium from the crystals. This will be discussed in detail in the loading capacity section.

When 10 percent bone char was added to the system, the magnesium concentration in the liquid phase doubled (Table A-7b). Calcium was strongly adsorbed from the groundwater at concentrations greater than 1.75 percent, and sodium was released into solution at concentrations greater than 1 percent. On a molar basis the amount of sodium dissolved does not match the amount of calcium adsorbed.

The hypothesis of protonation can be tested by analyzing changes in the pH condition of the liquid phase. Exchange of sodium for hydrogen should result in an increased hydroxyl abundance or a higher pH value. The pH for systems containing 20 percent bone char and clinoptilolite becomes 9.5 and 9.17, respectively (Table A-3); whereas, the same amount of NC apatite and HAP have little effect on the pH (8.0 and 8.2 respectively).

The concentration of anionic solutes is also influenced differently by the different materials. Figure 4-4 shows the elements sulfate, chlorine, and fluorine, normalized based on their molar abundances and plotted on a ternary diagram. Nitrate was excluded from the calculations, because its concentration was essentially constant under all experiments (Tables A-6a through A-6e). No determination of bicarbonate or total inorganic carbonate was made. Because carbonate is the major anion in the water, knowing its abundance could assist in the interpretation of the anion behavior.

Chlorine is strongly enriched relative to fluorine and sulfate in the solution when bone char is added. This would support the interpretation that this material may contain small amounts of halite ( $\text{NaCl}$ ) or sylvanite ( $\text{MgCl}_2$ ) that are simply dissolved when brought into contact with water. Ashing of animal bone and tissue would certainly be a plausible source of soluble alkali salts.

NC apatite has a tendency to increase the fluorine concentration of the groundwater (Figure 4-4). This can be explained by the re-equilibration of fluorapatite with the fluid and dissolving small amounts of the solid. Hydroxyapatite contributed only chlorine to the water because this synthetic mineral does not contain appreciable amounts of fluorine.

With the exception of NC apatite, all the materials that were tested adsorbed  $^{90}\text{Sr}$  from the solutions (Table A-5). NC apatite has a very high initial strontium content of  $1720 \mu\text{g/g}$  as determined by instrumental neutron activation analysis (INAA) and electron microprobe spectrometry (EMPS). In order to illustrate the sorption characteristics of the earth and alkali metals, the mass excess of sodium, magnesium, calcium, strontium, and barium was plotted against the weight fraction of HAP as a representative mineral for apatite in Figure 4-5. With the exception of strontium, all alkali and alkali earth elements show a mass deficit instead of a surface adsorption. For the lighter elements (Na, Mg, Ca) this is an indication of the ion exchange capacity of HAP. Barium concentrations did not vary significantly and the abundance of this element in the solutions was close to its detection limit. These factors are the main

contributors to the scatter shown in Figure 4-5. The mass excess of strontium follows a Langmuir isotherm on the graph, but at very low weight fractions of HAP the mass excess cannot be determined very accurately. This may be the result of the inherited sample inhomogeneity of the sediment sample. Only at very high concentrations of the homogeneous sorbent does this effect becomes less apparent. Nevertheless, the data indicate that strontium is adsorbed on the surfaces of hydroxyapatite.

The transition metals iron and chromium were desorbed from the system when hydroxyapatite was added (Table A-8c). Manganese showed a behavior different from the other metals. This element apparently will partition in the solid phases when phosphate is added. However, the results of the desorption and preliminary adsorption experiment indicated that manganese is highly reactive between 1 and 7 days. This could be a surface adsorption reaction or a secondary precipitation of manganese hydroxide that is linked to redox reactions in the system. Bone char has a similar effect on the manganese concentrations in the solution. The desorption and subsequent precipitation of metals from the placement of in situ barriers like bone char and clinoptilolite may potentially impact the hydraulic conductivity. Further investigation of this scenario is warranted.

The analytical data for zinc, chromium, and copper in NC apatite (Table A-8a) is fairly variable and often at the limit of detection. Therefore the interpretation of these values is very limited.

The question of the actual form in which  $^{90}\text{Sr}$  is present in the soils is not a subject of this study. It can be assumed that much of the strontium occurs as surface adsorbed species on the mineral grains in the soil. Diffusion of strontium into the crystal lattices is unlikely under the thermodynamic conditions of the sediment. Strontium may also be present in the form of small strontium carbonate or strontium sulfate precipitates in the soil because the solubility of these minerals is 1.1 mg/L and 1.13 mg/L in cold water, and  $\text{HCO}_3^{2-}$  is the major anion in the groundwater. Therefore it is to be expected that at lower pH conditions, the concentration of strontium should increase.

#### 4.4 KINETICS OF COMPLEXATION

To test the efficiency of adsorption of the sorbent materials, the amount of  $^{90}\text{Sr}$  remaining in solution after the addition of sorbents was examined at various time intervals. The time intervals for contact time were 1, 3, 5, 10, 15, and 24 hours. The amount of  $^{90}\text{Sr}$  remaining in solution at each time interval was plotted relative to the control as sorption efficiency. The concentration of  $^{90}\text{Sr}$  attributed to the desorption of  $^{90}\text{Sr}$  from the soil/sorbent mixture was subtracted from the amount that would be expected in the solution if only soil were present in the system, then these data were plotted as a function of time. The data for  $^{90}\text{Sr}$  are presented in Table B-1. These data are presented graphically in Figure 4-6.

Of the materials tested, the clinoptilolite was the most efficient in reducing the solution concentration of  $^{90}\text{Sr}$ . At 10 hours, the clinoptilolite has reduced the 13,683 pCi/L  $^{90}\text{Sr}$  solution concentration by 88 percent. Bone char reduced the  $^{90}\text{Sr}$  concentration by 52 percent. Both

clinoptilolite and bone char illustrate a plateau after the tenth hour of contact, suggesting that maximum adsorption is occurring within the first 10 hours. The addition of NC apatite did not change the desorption rate of  $^{90}\text{Sr}$  from the soil, which indicates that the natural apatite is not a suitable material in the stabilization of  $^{90}\text{Sr}$  in soils. However, when 5 percent hydroxyapatite or 5 percent bone char were added to the system, a significant reduction in the activity of  $^{90}\text{Sr}$  was observed. The kinetics of the complexation is fairly fast. It is noteworthy that the differential rate is similar for bone char and clinoptilolite.

#### 4.5 LOADING CAPACITY EXPERIMENT

The capacity of the sorbent material was measured by adding incremental amounts of strontium nitrate,  $\text{Sr}(\text{NO}_3)_2$ , to 3 g of NC apatite, bone char, hydroxyapatite, and clinoptilolite. The contact time for all three concentrations of  $\text{Sr}(\text{NO}_3)_2$  was 48 hours. The distribution coefficient ( $K_d$ ) was then calculated as the ratio of the strontium concentration adsorbed on the solid phase versus the strontium concentration remaining in the solution. The partition coefficients were then plotted in Figure 4-7. Data generated by this experiment are contained in Table C-1. At low strontium concentrations in the solutions, the cation is strongly adsorbed on most of the minerals. As the abundance of the solute increases, the capacity of the adsorbent decreases. A  $K_d$  of less than 1 indicates that more strontium remained in solution than was adsorbed on the crystal surfaces. The  $K_d$  values are substantially higher for the clinoptilolite and bone char for all concentrations of the  $\text{Sr}(\text{NO}_3)_2$ . The clinoptilolite has slightly higher  $K_d$  values than bone char, ranging from 585 at low concentrations of  $\text{Sr}(\text{NO}_3)_2$  to 0.41 for high concentrations of  $\text{Sr}(\text{NO}_3)_2$ . The bone char  $K_d$ s range from 371 at low concentrations of  $\text{Sr}(\text{NO}_3)_2$  to 0.31 for high concentrations of  $\text{Sr}(\text{NO}_3)_2$ .

Differences in  $K_d$ s may be substantially affected by surface area of the sorbents. The bone char, supplied by the Tigg Corporation, is reported to have a total surface area of  $100 \text{ m}^2/\text{g}$ . The clinoptilolite, supplied by the American Resource Corporation, Inc., is reported to have a total surface area of  $40 \text{ m}^2/\text{g}$ . Upon initial observation, it would appear that clinoptilolite is the better sorbent of the two, given the fact that the clinoptilolite has substantially lower surface area than bone char. It must be noted that the clinoptilolite and the bone char sorb  $^{90}\text{Sr}$  by two different chemical processes. The clinoptilolite is an ion exchanger and as such sorbs  $^{90}\text{Sr}$  onto the exchange sites initially occupied by monovalent and divalent cations. The ion exchange process occurs very quickly for the clinoptilolite, as evidenced by the kinetic experiment (Figure 4-6). The fact that ion exchange is occurring is evidenced by the high concentration of Na in the supernatant after the soil-clinoptilolite reaction with water. Na is the primary exchanger cation for the clinoptilolite. The bone char is an apatitic (calcium phosphate) mineral and sorbs  $^{90}\text{Sr}$  by incorporating the metal into its crystal formation. This is referred to as isomorphic substitution. Ca is replaced in the crystal structure by the  $^{90}\text{Sr}$ . This is evidenced by the elevated Ca levels in the supernatant of the bone char isotherms. The kinetics of the isomorphic substitution is somewhat slower than the ion exchange process, as evidenced by the slower rate of  $^{90}\text{Sr}$  sorption for bone char in Figure 4-6. The efficiencies and kinetics of the chemical reactions overshadow the apparent availability of reactive surface area when bone char is compared to clinoptilolite. Hydroxyapatite and NC apatite have less ability to sorb  $^{90}\text{Sr}$  than clinoptilolite and bone char.

#### 4.6 pH STABILITY TEST OF $^{90}\text{Sr}$ -PHOSPHATE COMPLEX

The stability of the  $^{90}\text{Sr}$ -phosphate complex was investigated under various pH conditions and reaction times as outlined in the scope of work. The soil was mixed with phosphate and clinoptilolite at proportions that were most effective for the stabilization of the contaminant. A total of 1 kg soil-sorbent mix was used for each experimental condition. Three replicates of each mixture were prepared by saturating with Hanford simulated groundwater and allowing the sample to dry to field capacity (0.33 bar). Each of the samples was then incubated for 3 different time periods: 1, 2, and 4 weeks. After incubation, an aliquot of the soil-sorbent mixture was removed and placed in a column. Six columns for each treatment were prepared to study the column water flow-through at pH levels of 5, 6, 7, 8, and 9. A control column that contained soil without added sorbent material was prepared for each of the treatments. A volume of Hanford simulated groundwater equal to 5 pore volumes of material and adjusted for the proper pH was introduced into the columns, and the reacted water was analyzed.

In the first phase of this experiment, a saturation extract was performed on 200 g of soil from each column. The soil was saturated with groundwater and the liquid phase was extracted and analyzed for  $^{90}\text{Sr}$ , ICP-MS metals, IC anions, pH and conductivity. The analytical results are presented in Tables D-1 through D-5. Although no control was performed for the saturation extract, the resultant data does indicate the immobilizing capability of the sorbents, especially given the incubation times of 1, 2, and 4 weeks. The percent reduction in solution  $^{90}\text{Sr}$  concentration is figured relative to NC apatite stabilization. The data for the resultant  $^{90}\text{Sr}$  concentration is plotted in column format and is shown in Figure 4-8. This figure indicates that both bone char- $^{90}\text{Sr}$  complex and clinoptilolite at 5-percent additions to the soil are remarkable in reducing the solution concentration of  $^{90}\text{Sr}$ . It must be noted that because of the small amount of water used in the saturation extract, the desorbed  $^{90}\text{Sr}$  is actually concentrated relative to  $^{90}\text{Sr}$  levels in the adsorption isotherm experiment. The clinoptilolite indicates a greater than 90 percent reduction over NC apatite stabilization, regardless of incubation time. The bone char sorbent is also exceptional in its ability to stabilize  $^{90}\text{Sr}$  with 90 percent reductions over NC apatite stabilization. Increasing the incubation time does not proportionately increase the stabilization factor for the bone char. Hydroxyapatite reduces  $^{90}\text{Sr}$  solution concentration more than the NC apatite, but is distinctly inferior to the capabilities of the bone char and clinoptilolite.

To test the stability of these sorbent complexes, 5 pore volumes each of pH levels 5, 6, 7, 8, and 9 flowed through 200 g columns taken from the 1 week, 2 week, and 4 week incubated soil. The column eluent was analyzed for  $^{90}\text{Sr}$  in solution. The data are presented in Table D-2 and graphically presented in Figure 4-9. The data are normalized against the control and represented as dissolution efficiency percent.

Both the clinoptilolite and bone char soil mixtures show excellent resistance to acidic and alkaline dissolution. It is quite surprising that little difference in eluent  $^{90}\text{Sr}$  concentrations exists between the different incubation times. The bone char indicates a slightly better stabilizing effect at lower pH with increasing incubation time. The bone char allows approximately 10 percent dissolution of the control concentration of  $^{90}\text{Sr}$  for the pH range of 4-9 (the control  $^{90}\text{Sr}$

concentration ranges from 15,489 pCi/L to 22,900 pCi/L). The clinoptilolite-<sup>90</sup>Sr complex resists dissolution 3 to 5 percent less than bone char, at pH levels of 5, 6, 7, 8, and 9.

The hydroxyapatite exhibits limited <sup>90</sup>Sr stability after incubation. The 4 week incubation soil/hydroxyapatite mixture proved the most resilient to dissolution, but only retained approximately 40 percent of the control solution <sup>90</sup>Sr. The resultant pH dissolution data for the NC apatite-<sup>90</sup>Sr complex was not presented in graph form as the amount of <sup>90</sup>Sr released by the varying pH solutions was greater than the control.

A general observation in this experiment is that the pH of the added groundwater solution was buffered by the soil after agitation. Whether either a mildly acidic solution at pH 5 or a basic solution was used, the pH after the groundwater was in contact for 1, 2, or 4 weeks ranged from pH 8 to 9. Therefore it would probably have been more beneficial for the interpretation of this experiment to counteract the natural buffering capacity of the soil by a buffered groundwater solution. This can be realized by adding, for instance, ammonium acetate and acetic acid to the groundwater solution. The amount of buffer should exceed the natural buffering capacity of the sediment.

Since the pH was buffered by the soil, it is not warranted to interpret the data as a function of pH in the groundwater solution.

## 4.7 SUMMARY OF RESULTS

The important results from each of the four experimental sections are summarized in the following sections.

### 4.7.1 Adsorption Isotherms

- Maximum desorption of <sup>90</sup>Sr from well 199-N-105A occurs within 72 hours. The concentration of <sup>90</sup>Sr at this time of desorption is approximately 12,000 pCi/L.
- The primary adsorption isotherm indicates that the clinoptilolite has the greatest ability to reduce <sup>90</sup>Sr solution concentration. The greatest reduction occurs at clinoptilolite values of less than 2 percent. The amount of <sup>90</sup>Sr precipitated and/or adsorbed at 2 percent added clinoptilolite is approximately 85 percent.
- The bone char is second in efficiency in reducing the <sup>90</sup>Sr concentration. All concentrations of bone char reduce <sup>90</sup>Sr concentration. However, 85 percent reduction in <sup>90</sup>Sr concentration is achieved at higher concentrations of bone char: between 10 and 20 percent. At 20 percent added bone char, the <sup>90</sup>Sr concentration is nearly the same as the reduction of <sup>90</sup>Sr caused by the addition of 2-percent clinoptilolite.

#### **4.7.2 Kinetics of Complexation**

- The clinoptilolite was most efficient in reducing the solution concentration of  $^{90}\text{Sr}$ .
- At 10 hours, the clinoptilolite reduced the 13,683 pCi/L  $^{90}\text{Sr}$  solution concentration by 88 percent.
- Bone char reduced the  $^{90}\text{Sr}$  concentration by 52 percent at the tenth hour of contact.
- Both clinoptilolite and bone char illustrate a plateau after the tenth hour of contact, suggesting that maximum adsorption is occurring within the first 10 hours.

#### **4.7.3 Loading Capacity Experiment**

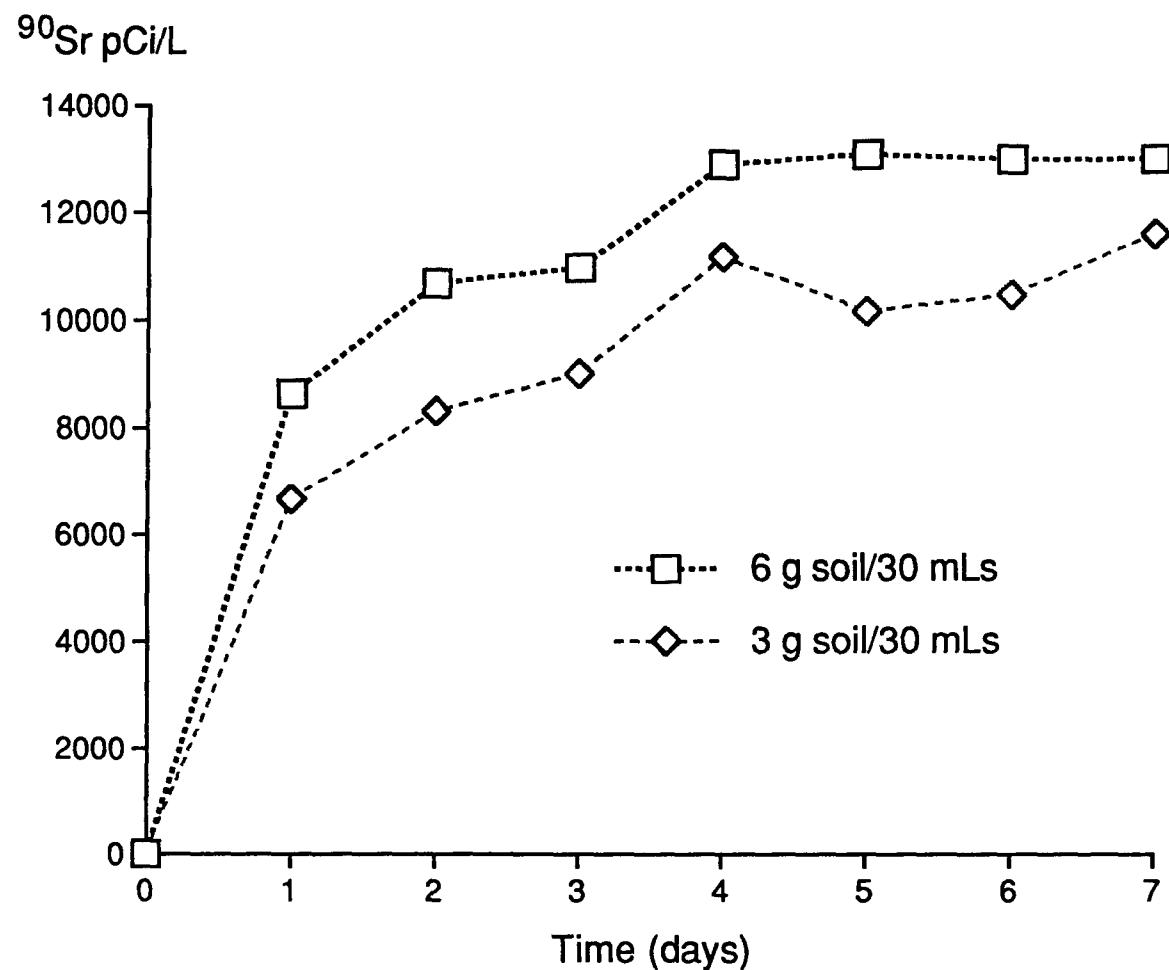
- The  $K_d$  values are highest for the clinoptilolite and bone char for all concentrations of the  $\text{Sr}(\text{NO}_3)_2$ .
- The clinoptilolite has slightly higher  $K_d$  values than bone char, ranging from 585 at low concentrations of  $\text{Sr}(\text{NO}_3)_2$  to 0.41 for high concentrations of  $\text{Sr}(\text{NO}_3)_2$ . The bone char  $K_d$ s range from 371 at low concentrations of  $\text{Sr}(\text{NO}_3)_2$  to 0.31 for high concentrations of  $\text{Sr}(\text{NO}_3)_2$ .

#### **4.7.4 pH Stability Test of $^{90}\text{Sr}$ -Phosphate Complex**

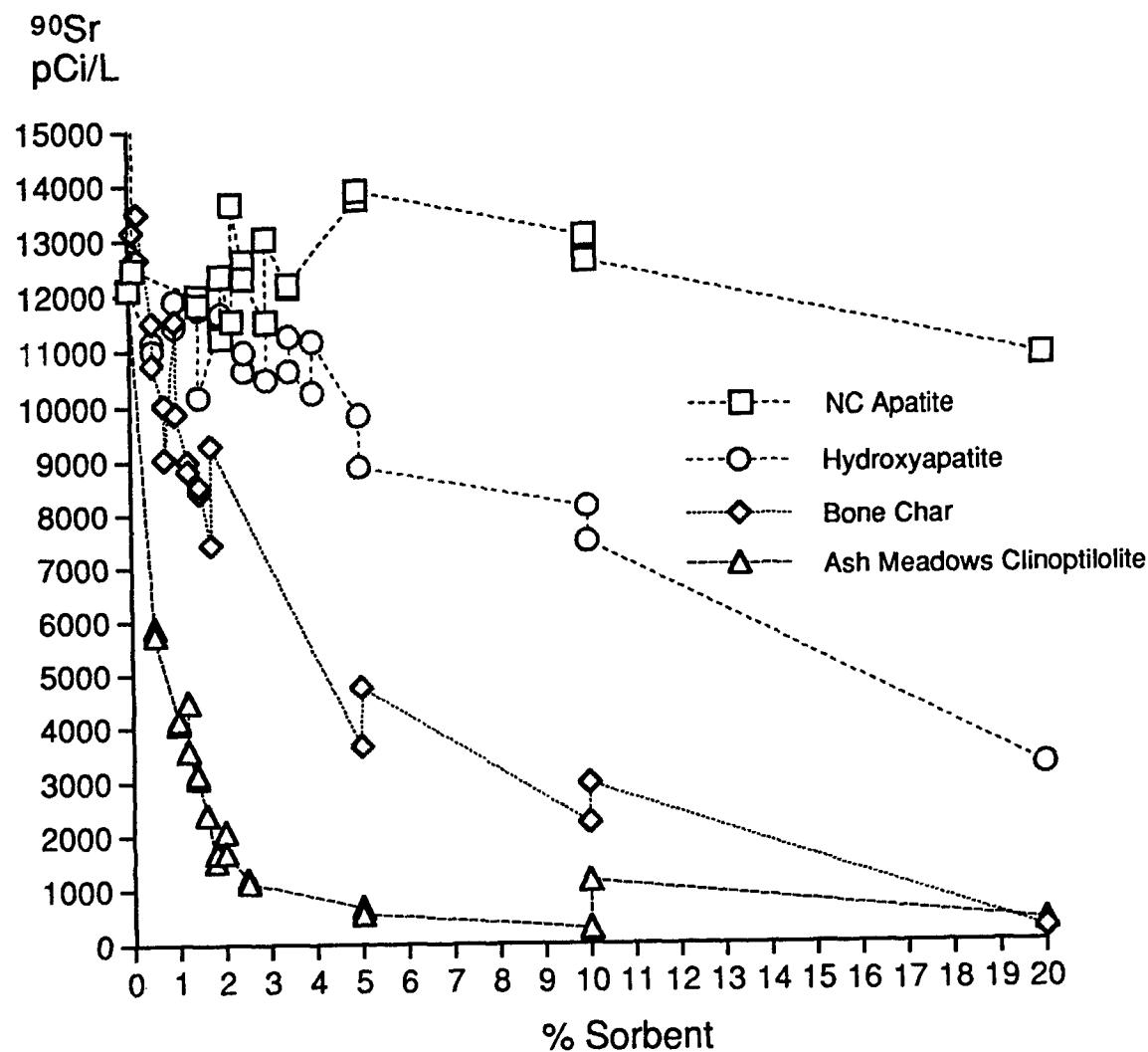
- Bone char and clinoptilolite at 5 percent additions to the soil are remarkable in reducing the solution concentration of  $^{90}\text{Sr}$ .
- The clinoptilolite indicates a greater than 90 percent reduction in solution concentration of  $^{90}\text{Sr}$  over NC apatite stabilization. This is the case whether at 1 week or 4 weeks of incubation time.
- The bone char sorbent is also exceptional in its ability to stabilize  $^{90}\text{Sr}$ , with 90 percent reductions in solution concentration of  $^{90}\text{Sr}$  over NC apatite stabilization.
- Increasing the incubation time does not proportionately increase the stabilization factor for the bone char.
- Clinoptilolite and bone char soil mixtures show excellent resistance to acidic and alkaline dissolution of the  $^{90}\text{Sr}$ -sorbent complex.
- The bone char allows approximately 10 percent of the control concentration of  $^{90}\text{Sr}$  for the pH range of 5 to 9 (the control  $^{90}\text{Sr}$  concentration ranges from 15,489 pCi/L to 22,900 pCi/L).

- The clinoptilolite is slightly better than the bone char, exhibiting less than 10 percent of the control concentration of pH levels of 5, 6, 7, 8, and 9.

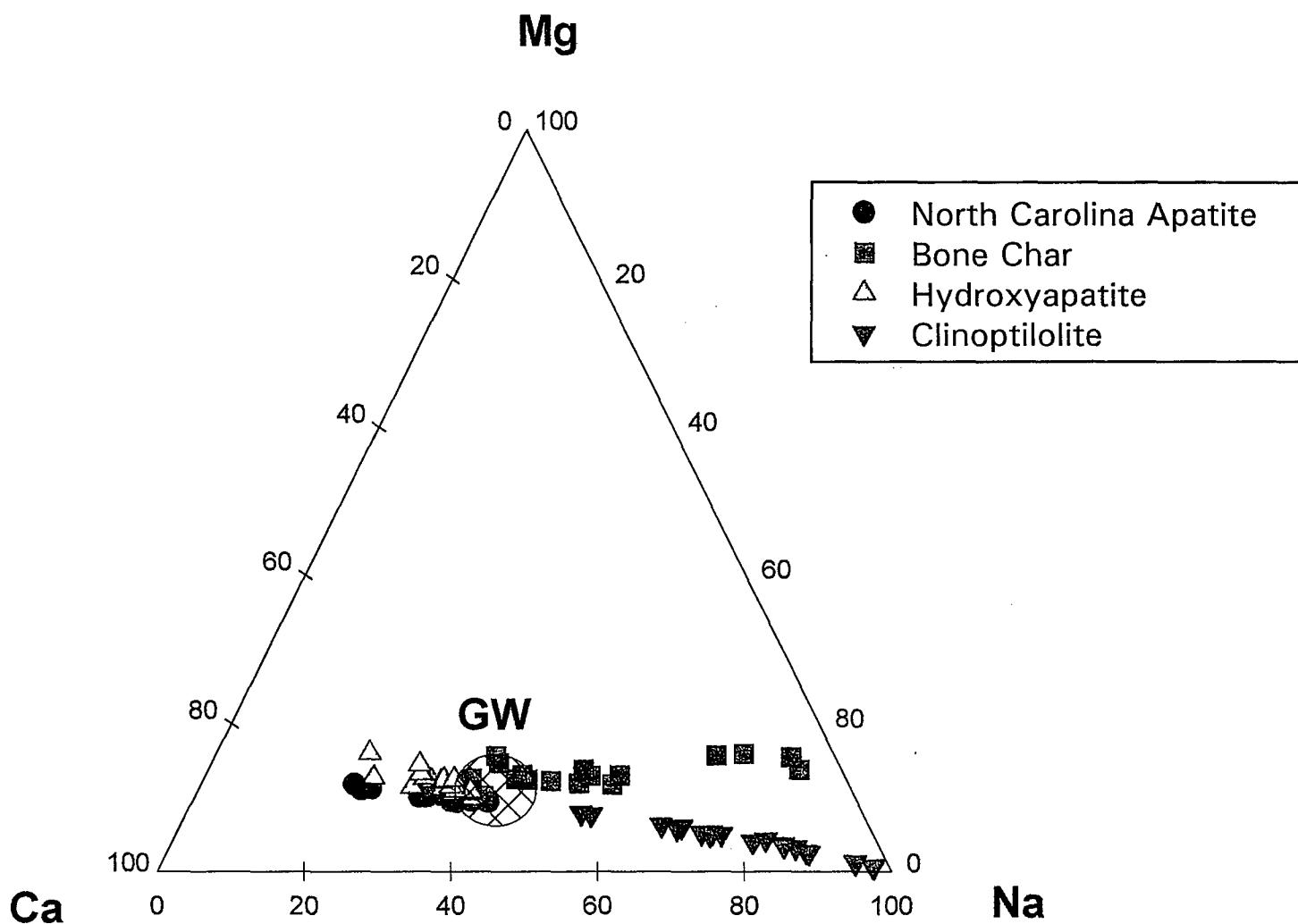
**Figure 4-1. Desorption of Strontium-90 from N-Springs Soil, Well 199-N-105A,  
Using Hanford Simulated Groundwater**



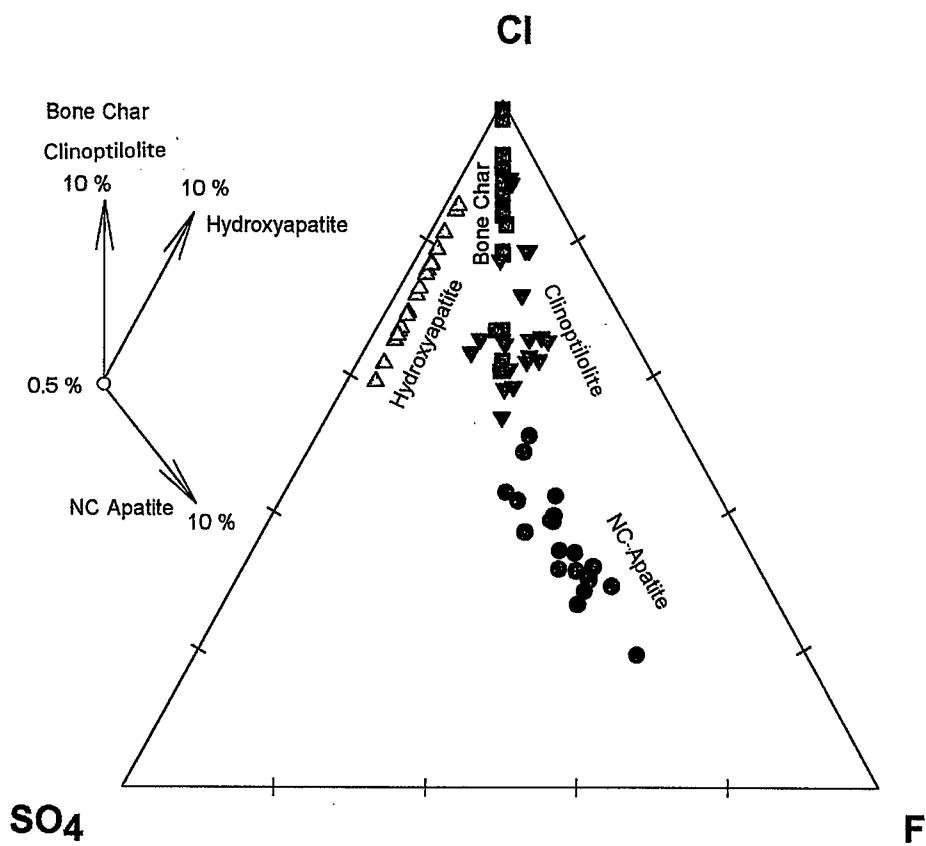
**Figure 4-2. Reduction of Solution  $^{90}\text{Sr}$  with the Addition of Incremental Amounts of Sorbent During 72 Hours Contact Time**



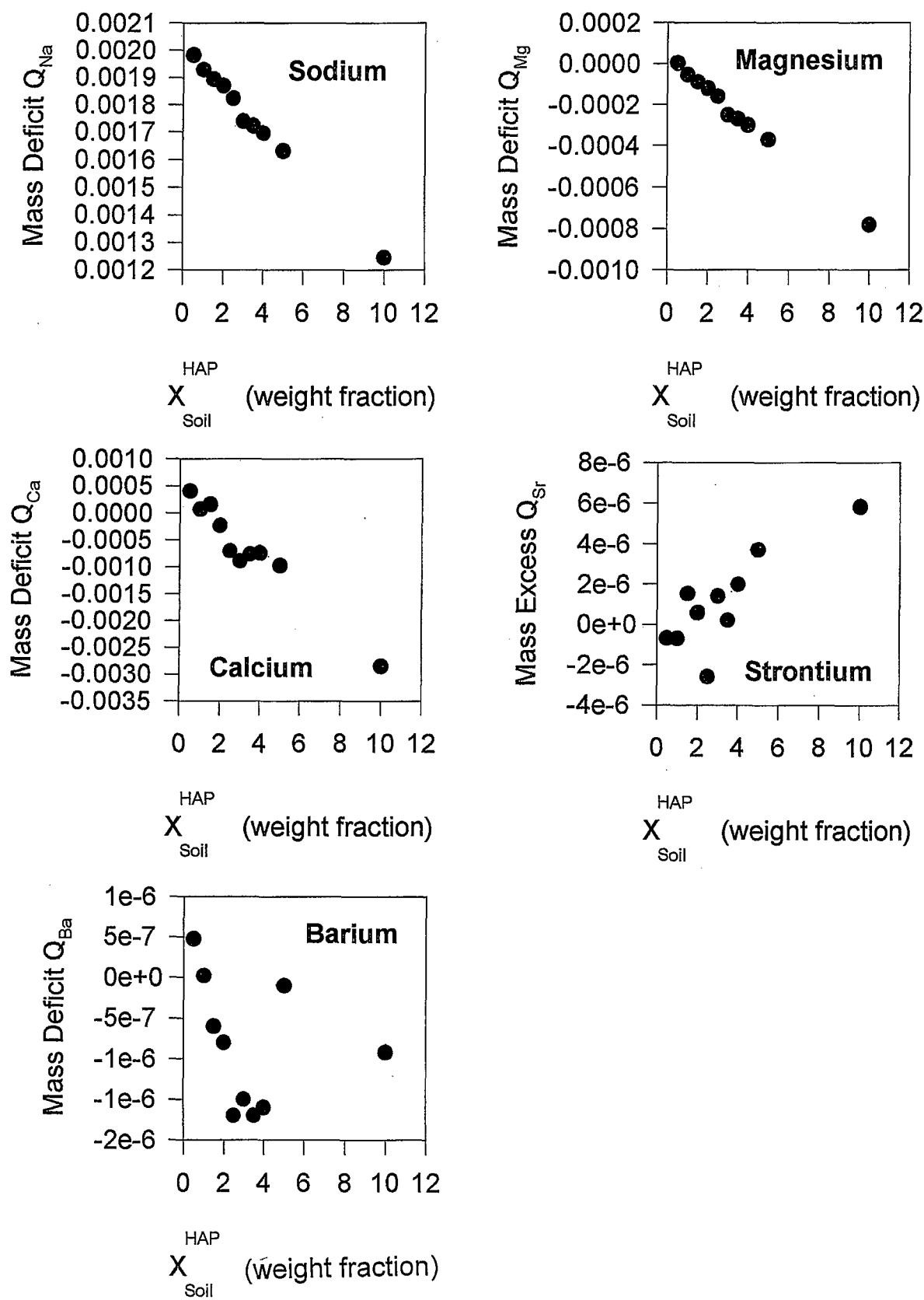
**Figure 4-3. Plot of Normalized Abundances of Major Element Compositions in Contact with Various Sorbent Materials**



**Figure 4-4. Changes in the Anion Composition of the Liquid Phase Illustrated by Normalized Element Concentrations**



**Figure 4-5. Plot of the Weight Fraction of HAP Versus the Mass Excess (Adsorbed Solute) for Alkali and Earth Alkali Elements**



**Figure 4-6. Time Required for Strontium-90 Sorption Using a 10:1 Water-to-Soil Ratio and a Fixed Amount of Sorbent Material**

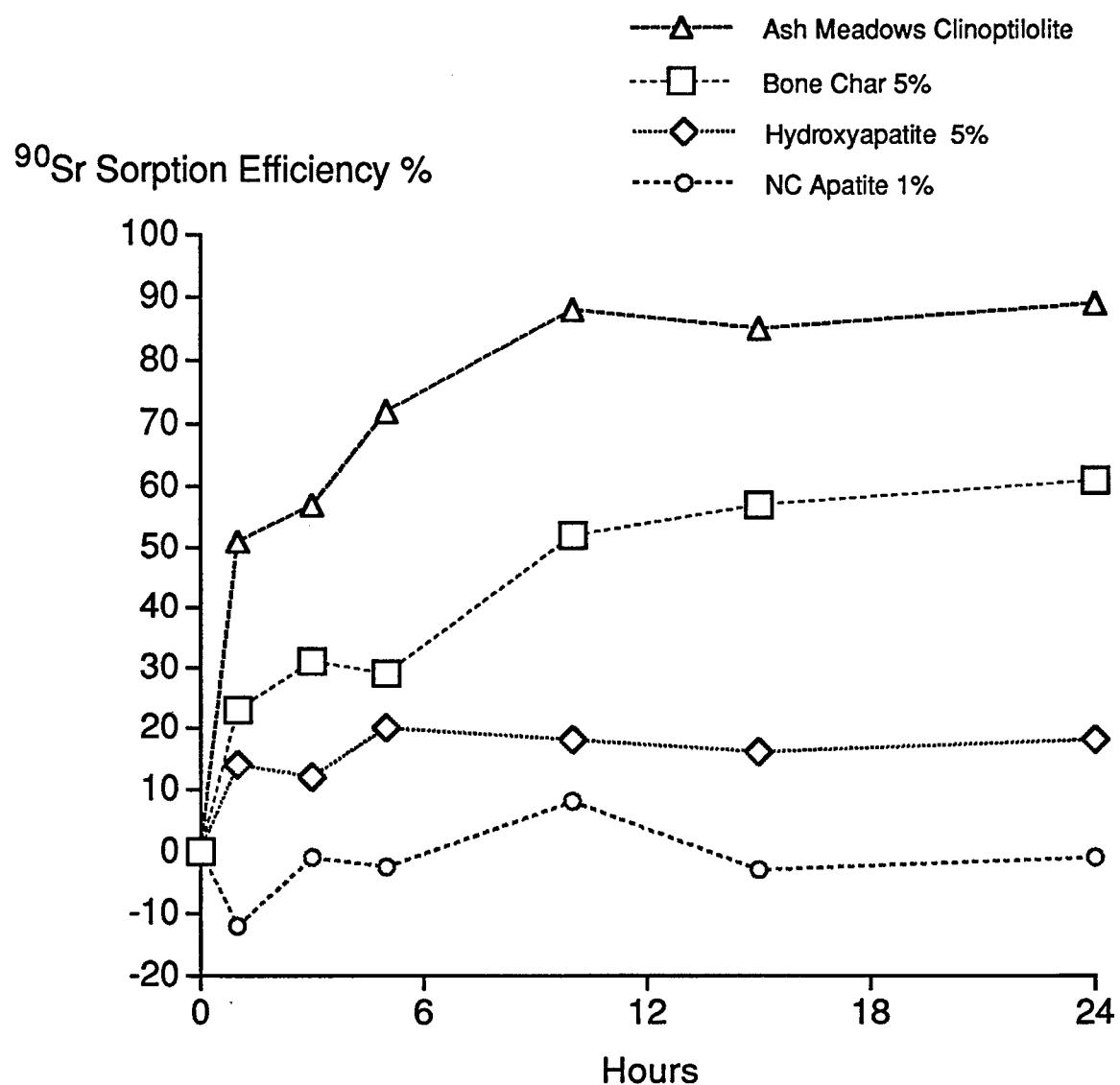


Figure 4-7. Strontium-90 Loading Capacity of the Sorbents Expressed as  $K_d$  Values

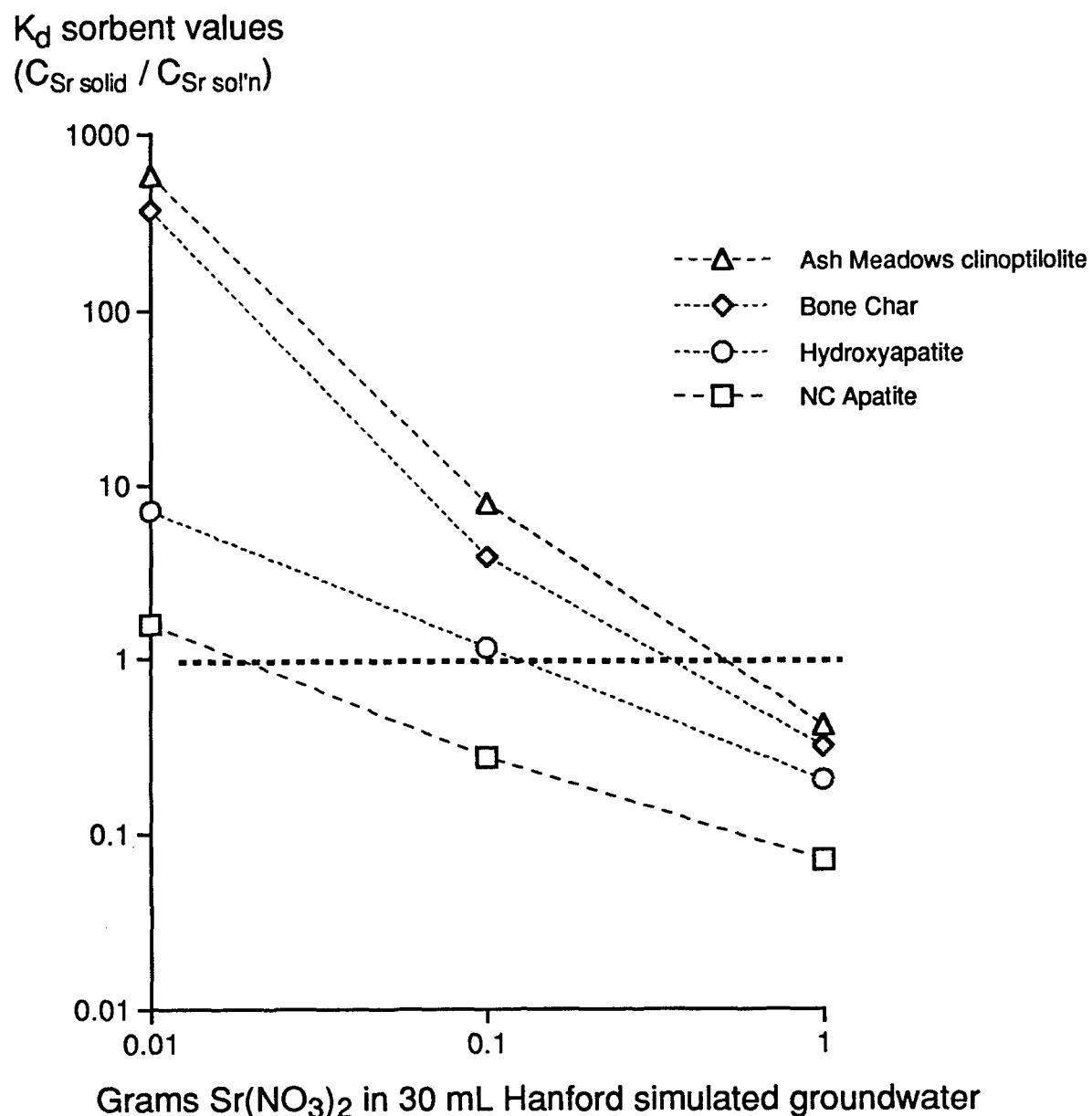


Figure 4-8. Strontium-90 in Solution After Complexed with Sorbent for 1, 2, and 4 Weeks

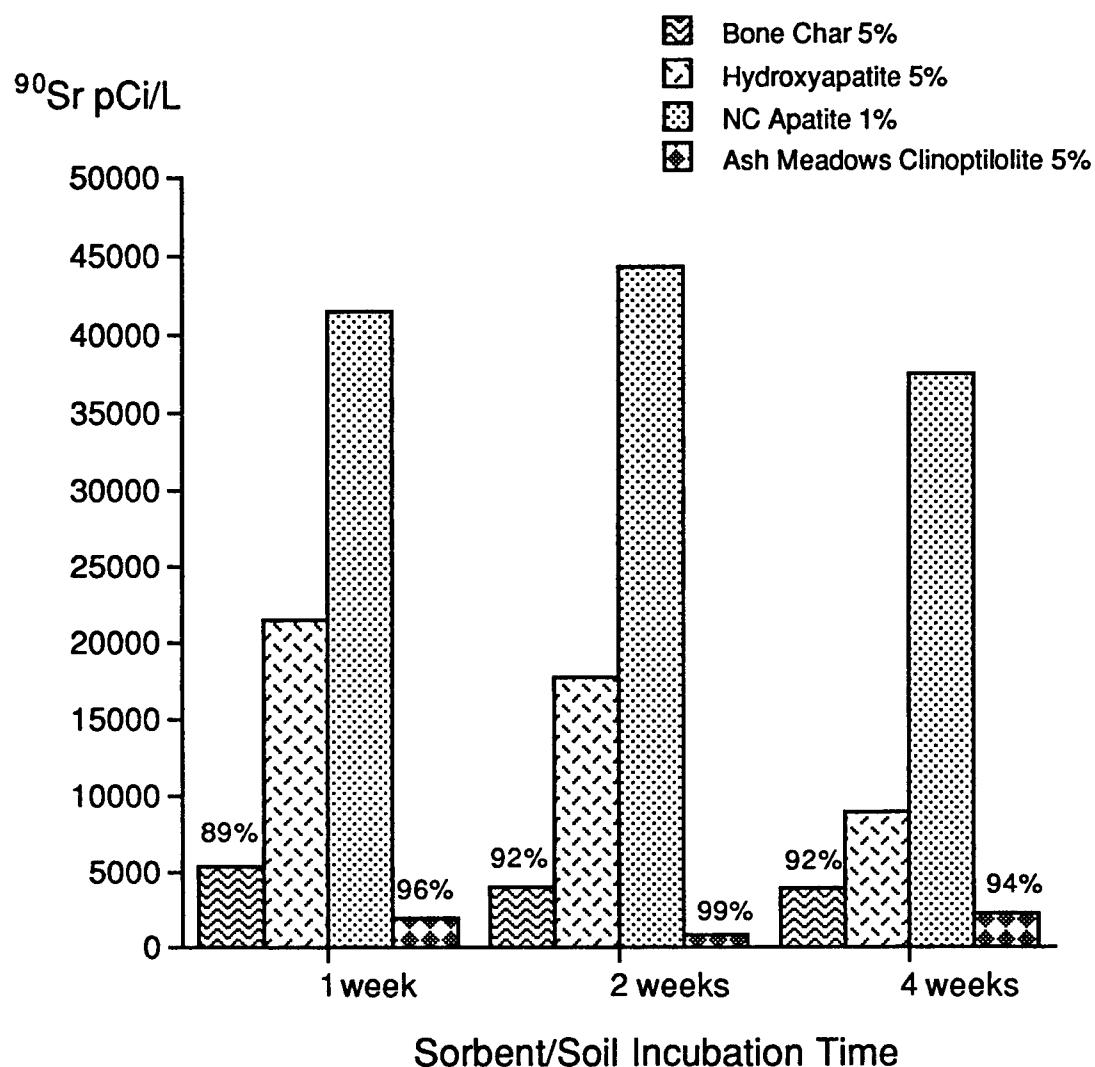
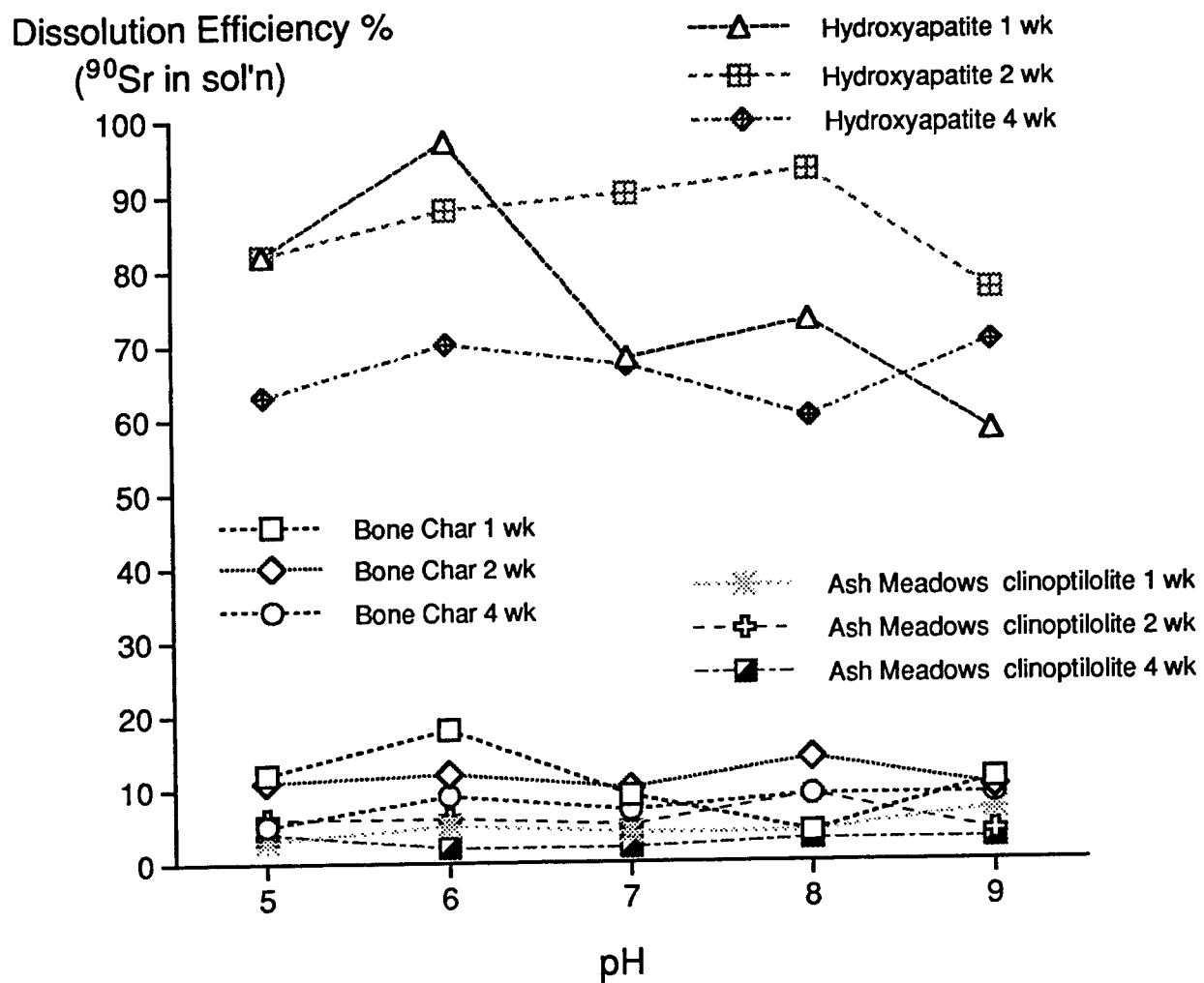


Figure 4-9. pH Stability of the Incubated 1, 2, and 4 Week  $^{90}\text{Sr}$ -Sorbent Complexes



BHI-00864  
Rev. 0

## 5.0 REFERENCES

- Akizuki, M., and Sunagawa, I., "Study of the Sector Structure in Adularia by Means of Optical Microscopy, Infra-Red Absorption, and Electron Microscopy." *Mineralogical Magazine*, 1994: 453-63.
- Anderson, A. J., "Influence of Phosphorus and Nitrogen Nutrition on Uptake and Distribution of Strontium and Calcium in Oat Plants." *Soil Science Society of America*, 1971: 308-11.
- Carter, J., 1990, *Biomineralization: Patterns, Process and Evolutionary Trends*, Van Nostrand Reinhold, New York, New York.
- Chandler, S., and Fuerstenau, D. W., 1984, "Solubility and Interfacial Properties of Hydroxyapatite: A Review." Misra, D. N., (Ed.), *Adsorption on and Surface Chemistry of Hydroxyapatite*, Plenum Press, New York, 177 pages.
- Deer, W. A., Howie, R. A., and Zussman, J., 1992, *An Introduction to the Rock-Forming Minerals*, Second Edition. Longman Scientific and Technical, Harlow, Essex, England.
- Edgar, A.D., "Barium and Strontium Enriched Apatites in Lamproites from West Kibreley, Western Australia." *American Mineralogist*, 1989: 889-95.
- Francis, C. W., "Radiostrontium Movement in Soils and Uptake by Plants." *National Technical Information Service*, 1978.
- Hughes, J. M., Cameron, M., Crowley, K. D., "Structural Variations in Natural F, OH, and Cl Apatites." *American Mineralogist*, 74 (1989): 870-76.
- Hughes, J. M., Cameron, M., Mariano, A. N., "Rare-Earth-Element Ordering and Structural Variations in Natural Rare-Earth-Bearing Apatites." *American Mineralogist*, 76 (1991) 1165-73.
- Liu, Y., and Comodi, P., "Some Aspects of the Crystal Chemistry of Apatites." *Mineralogical Magazine*, 1993: 709-19.
- Moody, T. E., and J. V. Wright, 1995, *Adsorption Isotherms: North Carolina Apatite Induced Precipitation of Lead, Zinc, Manganese, and Cadmium from the Bunker Hill 4000 Soil*, BHI-00197, Rev. 00, Bechtel Hanford Inc., Richland, Washington.
- Serne, R. J., and V. L. LeGore, 1996, *Strontium-90 Adsorption-Desorption Properties and Sediment Characterization at the 100 N-Area*, PNL-10899, Pacific Northwest Laboratory, Richland, Washington.
- Sposito, G., 1989, *The Chemistry of Soils*, Oxford University Press, New York, New York.

Rakován, J., and Reeder, R. J., "Differential Incorporation of Trace Elements and Dissymmetrization in Apatite: the Role of Surface Structure During Growth." *American Mineralogist*, 1994: 892-903.

Verbeeck, R.M. H., Hauben, M., Thun, H. P., and F. Verbeek, "Solubility and Solution Behaviour of Stontiumhydroxyapatite" *Z. Phys. Chem. (Wiesbaden)*, 108.2 (1977) 203-15.

**APPENDIX A.**

**Experiment A: Primary Absorption Experiment**

BHI-00864  
Rev. 0

**Table A-1. Liquid Scintillation Counting of Strontium-90, and pH and Conductivity Data from the Preliminary Desorption Experiment**

	Time of desorption	Date of counting	Sr-90 Bq/mL	Sr-90 pCi/L	pH	Conductivity microSi/cm
3 g of soil	1 day	10/10/95	0.27	7,244	8.47	na
3 g of soil	1 day	10/10/95	0.23	6,099	8.44	na
3 g of soil	2 days	10/10/95	0.32	8,776	8.38	na
3 g of soil	2 days	10/10/95	0.29	7,871	8.35	na
3 g of soil	3 days	10/10/95	0.31	8,249	8.48	na
3 g of soil	3 days	10/10/95	0.36	9,809	8.38	na
3 g of soil	4 days	10/10/95	0.47	12,575	8.29	140
3 g of soil	4 days	10/10/95	0.36	9,788	8.37	147
3 g of soil	5 days	10/10/95	0.36	9,766	8.32	146
3 g of soil	5 days	10/11/95	0.39	10,596	8.33	146
3 g of soil	6 days	10/11/95	0.38	10,362	8.46	151
3 g of soil	6 days	10/11/95	0.39	10,604	8.43	150
3 g of soil	7 days	10/11/95	0.44	11,936	7.88	155
3 g of soil	7 days	10/11/95	0.42	11,284	7.96	152
6 g of soil	1 day	10/11/95	0.32	8,534	8.47	na
6 g of soil	1 day	10/11/95	0.32	8,761	8.4	na
6 g of soil	2 days	10/11/95	0.37	9,990	8.4	na
6 g of soil	2 days	10/11/95	0.42	11,403	8.29	na
6 g of soil	3 days	10/11/95	0.37	9,984	8.41	na
6 g of soil	3 days	10/11/95	0.44	11,991	8.39	na
6 g of soil	4 days	10/11/95	0.43	11,657	8.37	150
6 g of soil	4 days	10/11/95	0.52	14,147	8.37	158
6 g of soil	5 days	10/11/95	0.50	13,489	8.15	159
6 g of soil	5 days	10/11/95	0.47	12,708	8.31	162
6 g of soil	6 days	10/11/95	0.47	12,694	8.3	157
6 g of soil	6 days	10/12/95	0.49	13,308	8.42	158
6 g of soil	7 days	10/12/95	0.51	13,850	7.96	161
6 g of soil	7 days	10/12/95	0.45	12,160	7.98	157
<b>Hanford Water</b>		10/13/95	0	-	8.2	122

**Table A-2. Ion Chromotography Data from the Preliminary Desorption Experiment**

	Time of desorption	Fluoride ppm	Chloride ppm	Nitrate ppm	Phosphate ppm	Sulfate ppm
3 g of soil	1 day	na	na	na	na	na
3 g of soil	1 day	na	na	na	na	na
3 g of soil	2 days	na	na	na	na	na
3 g of soil	2 days	na	na	na	na	na
3 g of soil	3 days	na	na	na	na	na
3 g of soil	3 days	na	na	na	na	na
3 g of soil	4 days	<0.1	1.67	9.17	<0.5	18.2
3 g of soil	4 days	<0.1	1.72	8.89	3.20	18.1
3 g of soil	5 days	<0.1	1.71	9.19	<0.5	18.2
3 g of soil	5 days	<0.1	1.58	9.71	<0.5	18.3
3 g of soil	6 days	<0.1	1.68	8.93	<0.5	18.3
3 g of soil	6 days	<0.1	1.63	9.43	<0.5	18.3
3 g of soil	7 days	na	na	na	na	na
3 g of soil	7 days	na	na	na	na	na
6 g of soil	1 day	na	na	na	na	na
6 g of soil	1 day	na	na	na	na	na
6 g of soil	2 days	na	na	na	na	na
6 g of soil	2 days	na	na	na	na	na
6 g of soil	3 days	na	na	na	na	na
6 g of soil	3 days	na	na	na	na	na
6 g of soil	4 days	na	na	na	na	na
6 g of soil	4 days	<0.1	1.91	8.77	5.58	18.0
6 g of soil	5 days	<0.1	1.76	9.36	<0.5	19.0
6 g of soil	5 days	<0.1	1.65	9.35	<0.5	18.8
6 g of soil	6 days	<0.1	1.97	9.05	2.02	19.0
6 g of soil	6 days	<0.1	1.66	9.49	<0.5	18.7
6 g of soil	7 days	<0.1	1.65	9.03	<0.5	18.8
6 g of soil	7 days	<0.1	3.39	18.71	<0.5	37.4
<b>Hanford Water</b>		<0.1	0.35	12.17	<0.5	16.6
<b>Maximal error of analysis in percent</b>		4.72	4.36	1.77	0.65	1.32

**Table A-3. Liquid Scintillation Counting of Strontium-90, and pH and Conductivity Data from the Preliminary Adsorption Experiment**

	Time of desorption	Date of counting	Sr-90 Bq/mL	Sr-90 pCi/L	pH	Conductivity microSi/cm
<b>Hanford Water</b> <b>2 days desorption</b>		10/13/95	0	-	8.2	122
		10/10/95	0.31	8,323	8.37	na
<b>NC Apatite</b>	1%	10/12/95	0.33	9,042	8.02	128
	5%	10/12/95	0.39	10,522	7.98	144
	10%	10/12/95	0.39	10,478	8	160
	20%	10/12/95	0.40	10,858	8.02	181
	pure				8.15	165
<b>Bone Char</b>	1%	10/12/95	0.12	3,242	8.43	212
	5%	10/12/95	0.22	5,881	8.43	157
	10%	10/12/95	0.28	7,473	8.17	123
	20%	10/12/95	0.009	232	9.54	338
	pure				10.23	386
<b>Hydroxyapatite</b>	1%	10/12/95	0.30	7,999	8.02	128
	5%	10/12/95	0.19	5,138	8.06	147
	10%	10/12/95	0.19	5,243	8.15	172
	20%	10/13/95	0.12	3,243	8.21	195
	pure				8.19	131
<b>Clinoptilolite</b>	1%	10/13/95	0.19	5,238	8.15	132
	5%	10/13/95	0.062	1,671	8.24	149
	10%	10/13/95	0.009	239	8.66	190
	20%	10/13/95	0.014	377	9.17	233
	pure				8.37	181
<b>Ca(H<sub>2</sub>PO<sub>4</sub>)<sub>2</sub>. H<sub>2</sub>O</b>	1%	10/13/95	0.92	24,942	6.26	529
	5%	10/13/95	1.55	41,930	6.07	1894
	10%	10/13/95	1.81	48,917	4.43	2940
	20%	10/13/95	1.98	53,618	3.51	6220
	pure				2.97	6830
<b>Phosphoric Acid</b>	1%	10/13/95	1.32	35,657	3.86	1044
	5%	10/13/95	1.81	49,007	2.43	3040
	10%	10/13/95	2.16	58,388	2.08	5840
	20%	10/13/95	2.49	67,215	1.82	10050
	pure				3.04	204

**Table A-4. Ion Chromotography Data from the Preliminary Adsorption Experiment**

		Fluoride ppm	Chloride ppm	Nitrate ppm	Phosphate ppm	Sulfate ppm
<b>Hanford Water</b>		<0.1	0.35	12.2	<0.5	16.6
<b>NC Apatite</b>	1%	0.23	0.75	16.5	<0.5	39.6
	5%	0.34	0.45	8.20	<0.5	25.9
	10%	0.46	0.46	7.30	<0.5	34.7
	20%	0.53	0.50	6.76	<0.5	46.3
	pure	0.58	0.57	8.39	<0.5	50.2
<b>Bone Char</b>	1%	<0.1	15.2	7.75	<0.5	19.3
	5%	<0.1	7.35	9.85	<0.5	18.3
	10%	<0.1	1.66	8.98	<0.5	17.2
	20%	<0.1	3.37	8.30	<0.5	24.2
	pure	<0.1	3.50	11.6	<0.5	24.3
<b>Hydroxyapatite</b>	1%	<0.1	0.47	8.23	<0.5	16.8
	5%	<0.1	0.61	7.62	<0.5	15.6
	10%	<0.1	0.78	6.55	<0.5	13.7
	20%	<0.1	1.13	3.79	0.56	11.2
	pure	<0.1	1.20	6.29	0.93	10.2
<b>Clinoptilolite</b>	1%	<0.1	0.61	8.06	<0.5	17.3
	5%	<0.1	1.38	8.20	<0.5	18.8
	10%	<0.1	2.66	7.96	<0.5	20.5
	20%	<0.1	5.19	6.64	<0.5	23.9
	pure	<0.1	4.96	12.2	<0.5	24.9
<b>Maximal error of analysis in percent</b>		4.72	4.36	1.77	0.65	1.32
<b>Ca(H<sub>2</sub>PO<sub>4</sub>)<sub>2</sub> • H<sub>2</sub>O</b>	1%	<0.1	1.41	6.50	na	17.0
	5%	na	na	na	na	17.4
	10%	na	na	1.55	na	17.6
	20%	na	na	na	na	17.6
	pure	na	na	na	na	na
<b>Phosphoric Acid</b>	1%	na	na	na	na	19.2
	5%	na	na	na	na	na
	10%	<0.1	na	na	na	79.0
	20%	na	na	na	na	na
	pure	na	na	na	na	na
<b>Maximal error of analysis in percent</b>		11.7	52.6	1.65	54.6	1.22

**Table A-5. Liquid Scintillation Counting of Strontium-90, pH, and Conductivity Data from the Primary Adsorption Isotherm Experiment (1 of 2)**

Material Tested	Percent Material Added	Date of counting	Sr-90			Conductivity microSi/cm
			Bq/mL	pCi/L	pH	
NC Apatite	0.5 %	11/27/95	0.51	13780	9.36	136
	0.5 %	11/27/95	0.45	12149	8.76	134
	1 %	11/27/95	0.88	23829	8.50	131
	1 %	11/27/95	0.46	12472	8.34	139
	1.5 %	11/27/95	0.44	11981	8.29	138
	1.5 %	11/27/95	0.44	11824	8.46	129
	2 %	11/28/95	0.42	11235	8.51	132
	2 %	11/28/95	0.46	12342	8.48	136
	2.25 %	11/28/95	0.43	11531	8.44	133
	2.25 %	11/28/95	0.51	13650	8.38	147
	2.5 %	11/28/95	0.47	12609	8.54	135
	2.5 %	11/28/95	0.46	12299	7.72	137
	3 %	11/28/95	0.43	11515	9.56	135
	3 %	11/28/95	0.48	13019	9.36	132
	3.5 %	11/28/95	0.45	12136	8.71	132
	3.5 %	11/28/95	0.45	12179	8.40	130
	5 %	11/28/95	0.51	13733	8.13	165
Bone Char	5 %	11/28/95	0.51	13882	8.20	160
	10 %	11/28/95	0.48	13048	8.20	180
	10 %	11/28/95	0.47	12597	8.20	178
	0.1 %	11/28/95	0.47	12619	6.70	152
	0.1 %	11/28/95	0.49	13152	7.98	133
	0.2 %	11/29/95	0.47	12665	8.17	133
	0.2 %	11/29/95	0.50	13487	8.14	153
	0.5 %	11/29/95	0.43	11509	8.19	142
	0.5 %	11/29/95	0.40	10750	8.55	123
	0.75 %	11/29/95	0.37	10027	8.50	126
	0.75 %	11/29/95	0.33	9044	8.46	119
	1 %	11/29/95	0.43	11529	8.45	145
	1 %	11/29/95	0.36	9863	8.52	139
	1.25 %	11/29/95	0.33	8995	8.85	129
	1.25 %	11/29/95	0.33	8822	8.96	130
	1.5 %	11/29/95	0.31	8418	9.02	131
	1.5 %	11/29/95	0.32	8539	9.08	132
	1.75 %	11/29/95	0.28	7438	9.11	135
	1.75 %	11/29/95	0.34	9288	8.71	143
	5 %	11/29/95	0.13	3643	9.33	193
	5 %	11/29/95	0.18	4745	9.27	184
	10 %	11/29/95	0.08	2216	9.55	272
	10 %	11/29/95	0.11	2953	9.42	273

**Table A-5. Liquid Scintillation Counting of Strontium-90, pH, and Conductivity Data from the Primary Adsorption Isotherm Experiment (2 of 2)**

Material Tested	Percent Material Added	Date of counting	Sr-90			Conductivity microSi/cm
			Bq/mL	pCi/L	pH	
<b>Hydroxyapatite</b>	0.5 %	11/29/95	0.41	11146	8.42	133
	0.5 %	11/29/95	0.41	10999	8.48	130
	1 %	11/29/95	0.42	11423	8.43	134
	1 %	11/29/95	0.44	11891	8.31	137
	1.5 %	11/29/95	0.43	11747	8.36	135
	1.5 %	11/29/95	0.38	10167	8.35	136
	2 %	11/29/95	0.43	11540	8.44	139
	2 %	11/29/95	0.43	11663	8.38	139
	2.5 %	11/29/95	0.39	10631	8.31	142
	2.5 %	11/29/95	0.41	10969	8.34	144
	3 %	11/29/95	0.39	10473	7.34	138
	3 %	11/29/95	0.43	11503	8.04	150
	3.5 %	11/29/95	0.42	11249	8.21	144
	3.5 %	11/29/95	0.39	10627	8.10	151
	4 %	11/29/95	0.38	10230	8.20	152
	4 %	11/29/95	0.41	11150	8.27	149
	5 %	11/29/95	0.36	9816	8.21	151
	5 %	11/29/95	0.33	8876	8.25	153
<b>Clinoptilolite</b>	10 %	11/29/95	0.30	8123	8.12	179
	10 %	11/29/95	0.28	7467	8.09	180
<b>Clinoptilolite</b>	0.5 %	11/29/95	0.22	5880	8.51	126
	0.5 %	11/29/95	0.21	5749	8.67	123
	1 %	11/29/95	0.15	4072	8.68	132
	1 %	11/29/95	0.15	4138	8.71	130
	1.2 %	11/30/95	0.17	4477	8.64	131
	1.2 %	11/30/95	0.13	3569	8.81	135
	1.4 %	11/30/95	0.11	3074	8.63	140
	1.4 %	11/30/95	0.12	3143	8.71	134
	1.6 %	11/30/95	0.09	2395	8.72	139
	1.6 %	11/30/95	not detected		8.71	135
	1.8 %	11/30/95	0.06	1527	8.89	141
	1.8 %	11/30/95	0.06	1700	8.89	137
	2 %	11/30/95	0.08	2070	8.94	142
	2 %	11/30/95	0.06	1682	8.90	132
	2.5 %	11/30/95	0.04	1202	9.12	150
	2.5 %	11/30/95	0.04	1133	9.03	148
	5 %	11/30/95	0.02	663	9.15	177
	5 %	11/30/95	0.02	550	9.22	172
<b>Blank</b>	10 %	11/30/95	0.01	275	9.64	224
	10 %	11/30/95	0.04	1152	9.66	227
<b>Blank</b>	Hanford Water	12/8/95	0.48	12851	8.33	129
	Hanford Water				8.44	111

**Table A-6a. Ion Chromotography Data from the Primary Adsorption Isotherm Experiment: North Carolina Apatite**

Weight % of NC Apatite	Fluoride mg/L			Chloride mg/L			Nitrate mg/L			Phosphate mg/L			Sulfate mg/L		
0.5 %	0.16	±	0.01	0.25	±	0.02	11.9	±	0.27	0.53	±	0.01	20.2	±	0.34
0.5 %	0.13	±	0.01	0.16	±	0.01	11.9	±	0.27	<0.5			20.6	±	0.34
1 %	0.54	±	0.04	0.23	±	0.02	12.1	±	0.27	<0.5			21.4	±	0.36
1 %	0.16	±	0.01	0.14	±	0.01	12.2	±	0.28	0.54	±	0.01	21.6	±	0.36
1.5 %	0.22	±	0.02	0.22	±	0.02	12.1	±	0.27	<0.5			22.5	±	0.38
1.5 %	0.16	±	0.01	0.17	±	0.01	12.4	±	0.28	0.53	±	0.01	22.5	±	0.38
2 %	0.22	±	0.02	0.16	±	0.01	12.4	±	0.28	<0.5			23.6	±	0.39
2 %	0.21	±	0.02	0.19	±	0.01	12.9	±	0.29	<0.5			23.4	±	0.39
2.25 %	0.26	±	0.02	0.18	±	0.01	12.9	±	0.29	<0.5			23.9	±	0.40
2.25 %	0.23	±	0.02	0.20	±	0.02	12.6	±	0.28	0.55	±	0.01	24.1	±	0.40
2.5 %	0.22	±	0.02	0.14	±	0.01	12.7	±	0.29	<0.5			24.0	±	0.40
2.5 %	0.23	±	0.02	0.20	±	0.02	21.0	±	0.47	0.51	±	0.01	24.9	±	0.42
3 %	0.31	±	0.02	0.19	±	0.01	12.7	±	0.29	<0.5			25.2	±	0.42
3 %	0.31	±	0.02	0.17	±	0.01	15.9	±	0.36	<0.5			25.7	±	0.43
3.5 %	0.28	±	0.02	0.17	±	0.01	13.0	±	0.29	<0.5			26.5	±	0.44
3.5 %	0.29	±	0.02	0.15	±	0.01	13.3	±	0.30	<0.5			25.8	±	0.43
5 %	0.33	±	0.03	0.16	±	0.01	12.7	±	0.29	<0.5			32.1	±	0.54
5 %	0.35	±	0.03	0.19	±	0.01	12.2	±	0.27	<0.5			29.7	±	0.50
10 %	0.65	±	0.05	0.18	±	0.01	13.1	±	0.30	<0.5			42.8	±	0.71
10 %	0.56	±	0.04	0.28	±	0.02	14.1	±	0.32	0.59	±	0.01	39.2	±	0.66
	mmol/L			mmol/L			mmol/L			mmol/L			mmol/L		
0.5 %	8.5	±	0.6	7.1	±	0.5	192	±	4	5.6	±	0.1	210	±	4
0.5 %	6.7	±	0.5	4.5	±	0.3	193	±	4	<5.3			214	±	4
1 %	28.5	±	2.1	6.5	±	0.5	195	±	4	<5.3			222	±	4
1 %	8.2	±	0.6	4.0	±	0.3	197	±	4	5.7	±	0.1	224	±	4
1.5 %	11.7	±	0.9	6.3	±	0.5	195	±	4	<5.3			235	±	4
1.5 %	8.2	±	0.6	4.9	±	0.4	200	±	5	5.5	±	0.1	234	±	4
2 %	11.8	±	0.9	4.6	±	0.3	199	±	4	<5.3			245	±	4
2 %	11.2	±	0.8	5.3	±	0.4	208	±	5	<5.3			244	±	4
2.25 %	13.6	±	1.0	5.0	±	0.4	208	±	5	<5.3			249	±	4
2.25 %	12.0	±	0.9	5.8	±	0.4	203	±	5	5.8	±	0.1	251	±	4
2.5 %	11.7	±	0.9	4.1	±	0.3	205	±	5	<5.3			250	±	4
2.5 %	12.2	±	0.9	5.7	±	0.4	338	±	8	5.4	±	0.1	259	±	4
3 %	16.5	±	1.2	5.2	±	0.4	204	±	5	<5.3			262	±	4
3 %	16.1	±	1.2	4.8	±	0.4	256	±	6	<5.3			268	±	4
3.5 %	14.9	±	1.1	4.9	±	0.4	210	±	5	<5.3			276	±	5
3.5 %	15.1	±	1.1	4.2	±	0.3	214	±	5	<5.3			269	±	4
5 %	17.6	±	1.3	4.6	±	0.3	204	±	5	<5.3			335	±	6
5 %	18.4	±	1.4	5.4	±	0.4	196	±	4	<5.3			309	±	5
10 %	34.4	±	2.6	5.2	±	0.4	212	±	5	<5.3			446	±	7
10 %	29.2	±	2.2	7.8	±	0.6	228	±	5	6.2	±	0.1	408	±	7

**Table A-6b. Ion Chromotography Data from the Primary Adsorption Isotherm Experiment: Bone Char**

Weight % of Bone Char	Fluoride mg/L	Chloride mg/L	Nitrate mg/L	Phosphate mg/L	Sulfate mg/L
0.1 %	0.11 ± 0.01	0.30 ± 0.02	13.8 ± 0.3	<0.5	19.4 ± 0.3
0.1 %	0.10	0.33 ± 0.02	12.5 ± 0.3	<0.5	19.6 ± 0.3
0.2 %	<0.1	0.40 ± 0.03	12.8 ± 0.3	<0.5	19.6 ± 0.3
0.2 %	0.11 ± 0.01	0.38 ± 0.03	12.9 ± 0.3	0.54 ± 0.01	19.9 ± 0.3
0.5 %	<0.1	0.72 ± 0.05	13.3 ± 0.3	<0.5	19.6 ± 0.3
0.5 %	<0.1	0.70 ± 0.05	12.5 ± 0.3	<0.5	19.3 ± 0.3
0.75 %	<0.1	1.0 ± 0.1	12.7 ± 0.3	<0.5	19.5 ± 0.3
0.75 %	0.13 ± 0.01	1.0 ± 0.1	12.7 ± 0.3	<0.5	19.6 ± 0.3
1 %	<0.1	1.2 ± 0.1	13.1 ± 0.3	0.52 ± 0.01	19.5 ± 0.3
1 %	<0.1	1.3 ± 0.1	13.5 ± 0.3	<0.5	19.5 ± 0.3
1.25 %	0.12 ± 0.01	1.7 ± 0.1	13.0 ± 0.3	<0.5	19.5 ± 0.3
1.25 %	<0.1	1.5 ± 0.1	13.0 ± 0.3	<0.5	19.6 ± 0.3
1.5 %	<0.1	1.9 ± 0.1	13.0 ± 0.3	<0.5	19.5 ± 0.3
1.5 %	<0.1	2.0 ± 0.2	13.1 ± 0.3	<0.5	19.6 ± 0.3
1.75 %	<0.1	2.4 ± 0.2	13.1 ± 0.3	<0.5	19.7 ± 0.3
1.75 %	<0.1	2.3 ± 0.2	12.9 ± 0.3	<0.5	19.7 ± 0.3
5 %	<0.1	8.4 ± 0.6	17.2 ± 0.4	<0.5	20.9 ± 0.3
5 %	<0.1	7.6 ± 0.6	13.1 ± 0.3	<0.5	20.6 ± 0.3
10 %	<0.1	16.7 ± 1.3	12.5 ± 0.3	<0.5	22.8 ± 0.4
10 %	<0.1	17.1 ± 1.3	12.6 ± 0.3	<0.5	22.8 ± 0.4
Bone Char	mmol/L	mmol/L	mmol/L	mmol/L	mmol/L
0.1 %	5.9 ± 0.4	8.4 ± 0.6	223 ± 5	<5.3	202 ± 3
0.1 %	<5.3	9.2 ± 0.7	201 ± 5	<5.3	204 ± 3
0.2 %	<5.3	11.2 ± 0.9	207 ± 5	<5.3	204 ± 3
0.2 %	5.6 ± 0.4	10.8 ± 0.8	208 ± 5	5.7 ± 0.1	207 ± 3
0.5 %	<5.3	20.3 ± 1.5	214 ± 5	<5.3	204 ± 3
0.5 %	<5.3	19.6 ± 1.5	201 ± 5	<5.3	201 ± 3
0.75 %	<5.3	28 ± 2	205 ± 5	<5.3	203 ± 3
0.75 %	6.9 ± 0.5	27 ± 2	206 ± 5	<5.3	204 ± 3
1 %	<5.3	34 ± 3	212 ± 5	5.4 ± 0.1	203 ± 3
1 %	<5.3	37 ± 3	218 ± 5	<5.3	203 ± 3
1.25 %	6.1 ± 0.5	48 ± 4	209 ± 5	<5.3	203 ± 3
1.25 %	<5.3	42 ± 3	209 ± 5	<5.3	204 ± 3
1.5 %	<5.3	55 ± 4	209 ± 5	<5.3	203 ± 3
1.5 %	<5.3	56 ± 4	212 ± 5	<5.3	204 ± 3
1.75 %	<5.3	68 ± 5	211 ± 5	<5.3	206 ± 3
1.75 %	<5.3	66 ± 5	208 ± 5	<5.3	205 ± 3
5 %	<5.3	237 ± 18	277 ± 6	<5.3	217 ± 4
5 %	<5.3	215 ± 16	212 ± 5	<5.3	214 ± 4
10 %	<5.3	471 ± 36	201 ± 5	<5.3	237 ± 4
10 %	<5.3	482 ± 36	203 ± 5	<5.3	237 ± 4

**Table A-6c. Ion Chromatography Data from the Primary Adsorption Isotherm Experiment: Hydroxyapatite**

Weight % of Hydroxyapatite	Fluoride mg/L	Chloride mg/L	Nitrate mg/L	Phosphate mg/L	Sulfate mg/L
0.5 %	<0.1	0.18 ± 0.01	12.6 ± 0.3	<0.5	19.4 ± 0.3
0.5 %	<0.1	0.16 ± 0.01	12.8 ± 0.3	<0.5	19.4 ± 0.3
1 %	<0.1	0.17 ± 0.01	12.7 ± 0.3	0.56 ± 0.01	19.3 ± 0.3
1 %	<0.1	0.21 ± 0.02	12.5 ± 0.3	0.59 ± 0.01	19.3 ± 0.3
1.5 %	<0.1	0.21 ± 0.02	12.2 ± 0.3	<0.5	18.8 ± 0.3
1.5 %	<0.1	0.20 ± 0.01	12.4 ± 0.3	<0.5	19.1 ± 0.3
2 %	<0.1	0.20 ± 0.02	12.3 ± 0.3	0.56 ± 0.01	19.1 ± 0.3
2 %	<0.1	0.33 ± 0.02	12.4 ± 0.3	0.55 ± 0.01	19.0 ± 0.3
2.5 %	<0.1	0.24 ± 0.02	12.1 ± 0.3	<0.5	18.8 ± 0.3
2.5 %	<0.1	0.23 ± 0.02	12.1 ± 0.3	0.53 ± 0.01	18.7 ± 0.3
3 %	<0.1	0.28 ± 0.02	11.8 ± 0.3	0.53 ± 0.01	18.5 ± 0.3
3 %	<0.1	0.38 ± 0.03	11.7 ± 0.3	<0.5	18.5 ± 0.3
3.5 %	<0.1	0.26 ± 0.02	11.7 ± 0.3	<0.5	18.4 ± 0.3
3.5 %	<0.1	0.28 ± 0.02	11.5 ± 0.3	0.54 ± 0.01	18.4 ± 0.3
4 %	<0.1	0.31 ± 0.02	11.6 ± 0.3	0.56 ± 0.01	18.3 ± 0.3
4 %	<0.1	0.44 ± 0.03	11.8 ± 0.3	0.56 ± 0.01	18.7 ± 0.3
5 %	<0.1	0.33 ± 0.02	11.2 ± 0.3	0.56 ± 0.01	17.8 ± 0.3
5 %	<0.1	0.32 ± 0.02	11.2 ± 0.3	0.56 ± 0.01	17.5 ± 0.3
10 %	<0.1	0.49 ± 0.04	10.0 ± 0.2	0.57 ± 0.01	16.1 ± 0.3
10 %	<0.1	0.50 ± 0.04	9.7 ± 0.2	0.59 ± 0.01	15.7 ± 0.3
	mmol/L	mmol/L	mmol/L	mmol/L	mmol/L
0.5 %	<5.3	5.1 ± 0.4	203 ± 5	<5.3	202 ± 3
0.5 %	<5.3	4.4 ± 0.3	206 ± 5	<5.3	202 ± 3
1 %	<5.3	4.9 ± 0.4	205 ± 5	5.9 ± 0.1	201 ± 3
1 %	<5.3	6.1 ± 0.5	201 ± 5	6.2 ± 0.1	201 ± 3
1.5 %	<5.3	6.0 ± 0.5	198 ± 4	<5.3	195 ± 3
1.5 %	<5.3	5.6 ± 0.4	200 ± 5	<5.3	199 ± 3
2 %	<5.3	5.8 ± 0.4	198 ± 4	5.9 ± 0.1	198 ± 3
2 %	<5.3	9.3 ± 0.7	200 ± 5	5.8 ± 0.1	198 ± 3
2.5 %	<5.3	6.7 ± 0.5	195 ± 4	<5.3	196 ± 3
2.5 %	<5.3	6.5 ± 0.5	196 ± 4	5.5 ± 0.1	195 ± 3
3 %	<5.3	7.9 ± 0.6	190 ± 4	5.6 ± 0.1	193 ± 3
3 %	<5.3	10.7 ± 0.8	189 ± 4	<5.3	193 ± 3
3.5 %	<5.3	7.4 ± 0.6	189 ± 4	<5.3	192 ± 3
3.5 %	<5.3	7.8 ± 0.6	185 ± 4	5.7 ± 0.1	191 ± 3
4 %	<5.3	8.6 ± 0.7	187 ± 4	5.9 ± 0.1	191 ± 3
4 %	<5.3	12.5 ± 0.9	191 ± 4	5.9 ± 0.1	195 ± 3
5 %	<5.3	9.3 ± 0.7	181 ± 4	5.8 ± 0.1	186 ± 3
5 %	<5.3	9.0 ± 0.7	180 ± 4	5.9 ± 0.1	182 ± 3
10 %	<5.3	13.7 ± 1.0	161 ± 4	6.0 ± 0.1	168 ± 3
10 %	<5.3	14.2 ± 1.1	156 ± 4	6.2 ± 0.1	163 ± 3

**Table A-6d. Ion Chromatography Data from the Primary Adsorption Isotherm Experiment: Clinoptilolite**

Weight % of clinoptilolite	Flouride mg/L	Chloride mg/L	Nitrate mg/L	Phosphate mg/L	Sulfate mg/L
0.5 %	0.12 ± 0.01	0.28 ± 0.02	12.4 ± 0.3	<0.5	19.8 ± 0.3
0.5 %	<0.1	0.23 ± 0.02	12.2 ± 0.3	<0.5	19.8 ± 0.3
1 %	0.13 ± 0.01	0.33 ± 0.03	12.5 ± 0.3	<0.5	20.0 ± 0.3
1 %	0.07 ± 0.01	0.29 ± 0.02	12.6 ± 0.3	<0.5	20.0 ± 0.3
1.2 %	0.14 ± 0.01	0.31 ± 0.02	12.6 ± 0.3	<0.5	20.3 ± 0.3
1.2 %	0.08 ± 0.01	0.33 ± 0.02	12.6 ± 0.3	<0.5	20.1 ± 0.3
1.4 %	0.13 ± 0.01	0.40 ± 0.03	12.5 ± 0.3	<0.5	21.1 ± 0.4
1.4 %	0.13 ± 0.01	0.39 ± 0.03	12.9 ± 0.3	<0.5	20.6 ± 0.3
1.6 %	0.17 ± 0.01	0.41 ± 0.03	12.8 ± 0.3	<0.5	20.5 ± 0.3
1.6 %	0.21 ± 0.02	0.46 ± 0.04	12.3 ± 0.3	<0.5	20.6 ± 0.3
1.8 %	0.23 ± 0.02	0.57 ± 0.04	12.5 ± 0.3	<0.5	20.6 ± 0.3
1.8 %	0.11 ± 0.01	0.68 ± 0.05	12.5 ± 0.3	<0.5	20.4 ± 0.3
2 %	0.19 ± 0.01	0.49 ± 0.04	12.9 ± 0.3	0.54 ± 0.01	20.5 ± 0.3
2 %	0.19 ± 0.01	0.46 ± 0.03	12.5 ± 0.3	14.39 ± 0.26	21.9 ± 0.4
2.5 %	0.18 ± 0.01	0.67 ± 0.05	12.6 ± 0.3	1.48 ± 0.03	20.9 ± 0.3
2.5 %	0.26 ± 0.02	0.61 ± 0.05	12.8 ± 0.3	1.55 ± 0.03	21.2 ± 0.4
5 %	0.26 ± 0.02	1.21 ± 0.09	13.1 ± 0.3	1.01 ± 0.02	21.9 ± 0.4
5 %	0.23 ± 0.02	1.11 ± 0.08	12.8 ± 0.3	0.70 ± 0.01	21.5 ± 0.4
10 %	0.21 ± 0.02	2.20 ± 0.17	13.4 ± 0.3	0.83 ± 0.02	23.1 ± 0.4
10 %	0.20 ± 0.01	2.29 ± 0.17	13.7 ± 0.3	0.90 ± 0.02	23.4 ± 0.4
	mmol/L	mmol/L	mmol/L	mmol/L	mmol/L
0.5 %	6.4 ± 0.5	7.9 ± 0.6	201 ± 5	<5.3	206 ± 3
0.5 %	<5.3	6.6 ± 0.5	197 ± 4	<5.3	206 ± 3
1 %	6.9 ± 0.5	9.4 ± 0.7	202 ± 5	<5.3	208 ± 3
1 %	3.9 ± 0.3	8.1 ± 0.6	203 ± 5	<5.3	208 ± 3
1.2 %	7.3 ± 0.5	8.7 ± 0.7	204 ± 5	<5.3	211 ± 4
1.2 %	4.4 ± 0.3	9.2 ± 0.7	204 ± 5	<5.3	209 ± 3
1.4 %	6.6 ± 0.5	11.3 ± 0.9	202 ± 5	<5.3	219 ± 4
1.4 %	6.7 ± 0.5	11.0 ± 0.8	208 ± 5	<5.3	215 ± 4
1.6 %	9.0 ± 0.7	11.5 ± 0.9	206 ± 5	<5.3	213 ± 4
1.6 %	10.9 ± 0.8	13.1 ± 1.0	199 ± 4	<5.3	215 ± 4
1.8 %	12.1 ± 0.9	16.1 ± 1.2	202 ± 5	<5.3	214 ± 4
1.8 %	6.0 ± 0.5	19.0 ± 1.4	202 ± 5	<5.3	212 ± 4
2 %	9.8 ± 0.7	13.9 ± 1.0	208 ± 5	5.7 ± 0.1	214 ± 4
2 %	10.1 ± 0.8	13.0 ± 1.0	202 ± 5	151.6 ± 2.7	228 ± 4
2.5 %	9.5 ± 0.7	18.8 ± 1.4	203 ± 5	15.5 ± 0.3	218 ± 4
2.5 %	13.7 ± 1.0	17.3 ± 1.3	207 ± 5	16.3 ± 0.3	221 ± 4
5 %	13.6 ± 1.0	34.1 ± 2.6	211 ± 5	10.6 ± 0.2	228 ± 4
5 %	12.1 ± 0.9	31.2 ± 2.4	206 ± 5	7.3 ± 0.1	224 ± 4
10 %	11.0 ± 0.8	62.1 ± 4.7	216 ± 5	8.7 ± 0.2	241 ± 4
10 %	10.4 ± 0.8	64.6 ± 4.9	221 ± 5	9.5 ± 0.2	244 ± 4

**Table A-6e. Ion Chromatography Data from the Primary Adsorption Isotherm Experiment: Hanford Soil**

Soil without sorbent	Flouride mg/L	Chloride mg/L	Nitrate mg/L	Phosphate mg/L	Sulfate mg/L
	mmol/L	mmol/L	mmol/L	mmol/L	mmol/L
100 %	0.13 ± 0.01	0.10	12.8 ± 0.5	0.7 ± 0.01	19.8 ± 0.6
100 %	0.10 ± 0.01	<0.1	13.1 ± 0.5	0.7 ± 0.01	19.9 ± 0.6
100 %	0.13 ± 0.01	<0.1	13.2 ± 0.5	<0.5	19.9 ± 0.6
100 %	0.11 ± 0.01	<0.1	13.1 ± 0.5	0.6 ± 0.01	20.0 ± 0.6
100 %	<0.1	<0.1	12.8 ± 0.5	<0.5	19.5 ± 0.6
100 %	6.6 ± 0.4	<2.8	207 ± 8	7.4 ± 0.1	206 ± 7
100 %	5.5 ± 0.3	<2.8	211 ± 8	7.1 ± 0.1	208 ± 7
100 %	6.7 ± 0.4	<2.8	213 ± 8	<5.3	207 ± 7
100 %	5.9 ± 0.3	<2.8	211 ± 8	6.3 ± 0.1	208 ± 7
100 %	<5.3	<2.8	206 ± 8	<5.3	203 ± 6

**Table A-7a. Atomic Absorption Spectrophotometry Data from the Primary Adsorption Isotherm Experiment: NC Apatite**

Weight % of NC Apatite	Sodium		Magnesium		Calcium	
	mg/L	StD%	mg/L	StD%	mg/L	StD%
0.5 %	7.1 ± 0.7	9.8	4.0 ± 0.02	0.5	22.8 ± 0.3	1.5
0.5 %	15.6 ± 0.2	1.2	3.7 ± 0.03	0.8	19.6 ± 0.4	2.1
1 %	13.7 ± 0.2	1.5	3.8 ± 0.001	0.03	19.8 ± 0.1	0.5
1 %	15.3 ± 0.2	1.5	4.0 ± 0.03	0.8	23.1 ± 0.3	1.1
1.5 %	14.4 ± 0.2	1.1	3.9 ± 0.01	0.2	22.5 ± 0.3	1.5
1.5 %	14.0 ± 0.3	1.9	3.7 ± 0.02	0.5	20.2 ± 0.7	3.5
2 %	13.6 ± 0.1	0.8	3.7 ± 0.01	0.4	19.7 ± 1.5	7.5
2 %	12.6 ± 0.3	2.5	3.9 ± 0.01	0.3	20.6 ± 0.5	2.7
2.25 %	14.7 ± 0.3	2.3	3.7 ± 0.01	0.2	20.3 ± 0.8	4.0
2.25 %	8.6 ± 1.6	18.2	4.1 ± 0.03	0.8	23.9 ± 0.3	1.4
2.5 %	16.4 ± 0.8	5.1	3.8 ± 0.03	0.8	20.4 ± 0.6	3.0
2.5 %	11.8 ± 0.0	0.0	3.8 ± 0.03	0.7	19.7 ± 1.9	9.4
3 %	13.5 ± 0.4	2.9	3.7 ± 0.02	0.6	19.1 ± 1.8	9.5
3 %	11.8 ± 0.4	3.8	3.7 ± 0.02	0.5	20.0 ± 2.2	10.8
3.5 %	13.1 ± 0.7	5.7	3.9 ± 0.04	1.2	20.8 ± 1.9	8.9
3.5 %	13.6 ± 0.7	5.4	3.8 ± 0.04	1.1	21.2 ± 3.0	14.3
5 %	13.3 ± 0.3	2.1	4.4 ± 0.03	0.7	25.7 ± 1.1	4.3
5 %	13.4 ± 0.1	0.7	4.3 ± 0.05	1.3	24.8 ± 1.0	4.1
10 %	14.6 ± 0.5	3.2	4.8 ± 0.02	0.5	27.7 ± 1.1	3.9
10 %	9.4 ± 0.5	5.6	4.7 ± 0.01	0.2	28.2 ± 1.2	4.3
NC Apatite	mmol/L	StD%	mmol/L	StD%	mmol/L	StD%
0.5 %	310 ± 30	9.8	165 ± 1	0.5	568 ± 8	1.5
0.5 %	679 ± 8	1.2	151 ± 1	0.8	488 ± 10	2.1
1 %	598 ± 9	1.5	155 ± 0	0.03	494 ± 2	0.5
1 %	667 ± 10	1.5	163 ± 1	0.8	576 ± 6	1.1
1.5 %	626 ± 7	1.1	162 ± 0	0.2	562 ± 9	1.5
1.5 %	609 ± 12	1.9	153 ± 1	0.5	503 ± 18	3.5
2 %	590 ± 5	0.8	153 ± 1	0.4	492 ± 37	7.5
2 %	546 ± 14	2.5	159 ± 0	0.3	514 ± 14	2.7
2.25 %	639 ± 15	2.3	154 ± 0	0.2	506 ± 20	4.0
2.25 %	373 ± 68	18.2	168 ± 1	0.8	597 ± 9	1.4
2.5 %	715 ± 36	5.1	155 ± 1	0.8	510 ± 15	3.0
2.5 %	513 ± 0	0.0	155 ± 1	0.7	490 ± 46	9.4
3 %	589 ± 17	2.9	152 ± 1	0.6	477 ± 45	9.5
3 %	514 ± 19	3.8	152 ± 1	0.5	500 ± 54	10.8
3.5 %	570 ± 33	5.7	159 ± 2	1.2	518 ± 46	8.9
3.5 %	590 ± 32	5.4	156 ± 2	1.1	528 ± 76	14.3
5 %	579 ± 12	2.1	183 ± 1	0.7	641 ± 28	4.3
5 %	583 ± 4	0.7	177 ± 2	1.3	620 ± 25	4.1
10 %	634 ± 20	3.2	196 ± 1	0.5	692 ± 27	3.9
10 %	407 ± 23	5.6	194 ± 0	0.2	704 ± 30	4.3

**Table A-7b. Atomic Absorption Spectrophotometry Data from the Primary Adsorption Isotherm Experiment: Bone Char**

Weight % of Bone char	Sodium mg/L	StD%	Magnesium mg/L	StD%	Calcium mg/L	StD%
0.1 %	15.0 ± 0.1	0.9	3.9 ± 0.01	0.2	19.2 ± 2.0	10.5
0.1 %	11.4 ± 0.2	1.6	4.0 ± 0.04	1.0	20.3 ± 2.0	9.8
0.2 %	14.2 ± 1.7	11.7	4.0 ± 0.001	0.03	19.6 ± 1.1	5.6
0.2 %	15.9 ± 0.1	0.8	4.6 ± 0.07	1.5	22.8 ± 1.1	4.7
0.5 %	13.7 ± 0.2	1.2	4.7 ± 0.06	1.3	19.1 ± 0.0	0.1
0.5 %	13.6 ± 1.2	9.1	4.0 ± 0.02	0.6	14.3 ± 0.3	2.1
0.75 %	14.7 ± 0.2	1.1	4.1 ± 0.05	1.1	14.4 ± 0.1	0.7
0.75 %	16.3 ± 0.5	3.1	4.2 ± 0.09	2.2	13.8 ± 0.4	2.8
1 %	16.9 ± 0.1	0.7	5.1 ± 0.03	0.7	17.1 ± 0.2	1.0
1 %	13.1 ± 1.2	8.9	4.9 ± 0.06	1.3	15.4 ± 0.2	1.1
1.25 %	11.1 ± 0.8	6.9	4.5 ± 0.06	1.4	13.3 ± 0.03	0.3
1.25 %	18.6 ± 0.1	0.3	4.3 ± 0.04	0.9	13.2 ± 0.2	1.8
1.5 %	21.5 ± 0.8	3.5	4.5 ± 0.01	0.3	12.3 ± 0.03	0.3
1.5 %	18.3 ± 0.7	4.0	4.5 ± 0.01	0.3	12.0 ± 0.6	4.7
1.75 %	20.4 ± 1.1	5.2	4.7 ± 0.01	0.2	11.0 ± 1.0	8.7
1.75 %	19.0 ± 0.7	3.6	5.1 ± 0.03	0.7	13.0 ± 1.4	10.7
5 %	34.5 ± 0.7	1.9	7.6 ± 0.02	0.3	5.8 ± 0.7	12.1
5 %	30.1 ± 0.3	0.9	6.9 ± 0.001	0.02	7.0 ± 0.7	9.7
10 %	53.4 ± 0.8	1.5	9.1 ± 0.01	0.1	3.8 ± 0.4	10.3
10 %	48.9 ± 0.9	1.8	9.6 ± 0.01	0.1	3.7 ± 0.3	7.4
Bone Char	mmol/L	StD%	mmol/L	StD%	mmol/L	StD%
0.1 %	651 ± 6	0.9	160 ± 0	0.2	479 ± 50	10.5
0.1 %	498 ± 8	1.6	163 ± 2	1.0	507 ± 49	9.8
0.2 %	619 ± 72	11.7	166 ± 0	0.03	488 ± 27	5.6
0.2 %	690 ± 6	0.8	187 ± 3	1.5	569 ± 27	4.7
0.5 %	596 ± 7	1.2	195 ± 3	1.3	477 ± 0	0.1
0.5 %	591 ± 54	9.1	165 ± 1	0.6	356 ± 7	2.1
0.75 %	639 ± 7	1.1	170 ± 2	1.1	360 ± 2	0.7
0.75 %	710 ± 22	3.1	174 ± 4	2.2	345 ± 10	2.8
1 %	736 ± 5	0.7	208 ± 1	0.7	427 ± 4	1.0
1 %	568 ± 50	8.9	200 ± 3	1.3	385 ± 4	1.1
1.25 %	485 ± 34	6.9	186 ± 3	1.4	331 ± 1	0.3
1.25 %	808 ± 3	0.3	178 ± 2	0.9	330 ± 6	1.8
1.5 %	934 ± 33	3.5	185 ± 1	0.3	307 ± 1	0.3
1.5 %	796 ± 31	4.0	186 ± 0	0.3	299 ± 14	4.7
1.75 %	889 ± 46	5.2	194 ± 0	0.2	275 ± 24	8.7
1.75 %	828 ± 29	3.6	209 ± 1	0.7	325 ± 35	10.7
5 %	1502 ± 29	1.9	314 ± 1	0.3	144 ± 17	12.1
5 %	1311 ± 12	0.9	283 ± 0	0.02	174 ± 17	9.7
10 %	2323 ± 36	1.5	376 ± 0	0.1	94 ± 10	10.3
10 %	2127 ± 38	1.8	397 ± 0	0.1	93 ± 7	7.4





**Table A-7e. Atomic Absorption Spectrophotometry Data from the Primary Adsorption Isotherm Experiment: Hanford Soil**

Hanford Soil	Sodium		Magnesium		Calcium	
	mg/L	StD%	mg/L	StD%	mg/L	StD%
100%	12.3 ± 0.3	2.5	3.7 ± 0.04	1.0	19.0	
100%	11.3 ± 1.0	9.0	3.6 ± 0.01	0.3	19.2	
100%	11.8 ± 0.3	2.7	3.8 ± 0.01	0.3	20.7	
100%	14.0 ± 0.0	0.3	3.8 ± 0.07	1.7	22.1	
100%	12.5 ± 0.0	0.3	4.5 ± 0.07	1.6	15.8 ± 0.4	2.8
100%	17.1 ± 0.4	2.3	4.5 ± 0.05	1.0	15.4 ± 0.2	1.4
<b>Average ±</b>	<b>13.2 ± 2.1</b>	<b>16.3</b>	<b>4.0 ± 0.4</b>	<b>9.6</b>	<b>18.7 ± 2.6</b>	<b>14.1</b>
<b>Standard deviation</b>						
	mmol/L	StD%	mmol/L	StD%	mmol/L	StD%
100%	0.54 ± 0.01	2.5	0.15 ± 0.001	1.0	0.47	
100%	0.49 ± 0.04	9.0	0.15 ± 0.000	0.3	0.48	
100%	0.51 ± 0.01	2.7	0.16 ± 0.001	0.3	0.52	
100%	0.61 ± 0.00	0.3	0.16 ± 0.003	1.7	0.55	
100%	0.54 ± 0.00	0.3	0.18 ± 0.003	1.6	0.39 ± 0.01	2.8
100%	0.75 ± 0.02	2.3	0.18 ± 0.002	1.0	0.39 ± 0.01	1.4
<b>Average ±</b>	<b>0.6 ± 0.1</b>	<b>16.3</b>	<b>0.16 ± 0.02</b>	<b>9.6</b>	<b>0.5 ± 0.1</b>	<b>14.1</b>
<b>Standard deviation</b>						

**Table A-8a. Inductively Coupled Plasma-Mass Spectrometry Data from the Primary Adsorption Isotherm Experiment: NC Apatite (1 of 4)**

Weight % of NC Apatite	Chromium µg/L	Manganese µg/L	Iron µg/L	Copper µg/L	Zinc µg/L
0.5 %	0.4 ± 0.0	7.5 ± 0.0	83 ± 2	4.2 ± 0.1	<1.4
0.5 %	0.3 ± 0.1	5.3 ± 0.0	73 ± 2	0.8 ± 0.0	1.4
1 %	0.3 ± 0.0	6.3 ± 0.1	83 ± 2	0.7 ± 0.0	<1.4
1 %	<0.2	8.4 ± 0.0	97 ± 3	<0.5	<1.4
1.5 %	<0.2	7.5 ± 0.0	61 ± 4	1.6 ± 0.0	2.6 ± 0.1
1.5 %	<0.2	5.6 ± 0.1	46 ± 1	1.0 ± 0.1	<1.4
2 %	<0.2	6.4 ± 0.1	54 ± 2	0.6 ± 0.0	<1.4
2 %	<0.2	7.2 ± 0.1	62 ± 3	1.1 ± 0.0	1.7 ± 0.2
2.25 %	<0.2	5.9 ± 0.1	45 ± 1	1.2 ± 0.1	<1.4
2.25 %	<0.2	9.1 ± 0.1	78 ± 2	1.2 ± 0.0	<1.4
2.5 %	<0.2	6.0 ± 0.1	51 ± 8	0.7 ± 0.0	<1.4
2.5 %	<0.2	7.1 ± 0.1	72 ± 4	0.6 ± 0.0	2.1 ± 0.2
3 %	0.3 ± 0.0	6.5 ± 0.0	57 ± 4	0.8 ± 0.0	2.0 ± 0.1
3 %	0.3 ± 0.0	6.8 ± 0.0	68 ± 4	0.8 ± 0.1	<1.4
3.5 %	0.2 ± 0.0	6.9 ± 0.1	52 ± 4	0.6 ± 0.0	<1.4
3.5 %	<0.2	6.3 ± 0.1	50 ± 3	0.6 ± 0.0	<1.4
5 %	<0.2	13.0	108	0.7	1.9
5 %	<0.2	10.7	88	<0.5	<1.4
10 %	<0.2	13.2 ± 0.0	118 ± 1	0.6 ± 0.0	<1.4
10 %	<0.2	14.6 ± 0.1	113 ± 3	1.1 ± 0.0	2.3 ± 0.0
NC Apatite	µmol/L	µmol/L	µmol/L	µmol/L	µmol/L
0.5 %	7.1 ± 0.4	136 ± 1	1492 ± 28	65 ± 1	<21
0.5 %	5.3 ± 1.0	96 ± 0	1299 ± 34	13 ± 1	<21
1 %	6.6 ± 0.9	115 ± 1	1493 ± 31	12 ± 0	<21
1 %	<3.8	153 ± 0	1734 ± 50	<8	<21
1.5 %	<3.8	136 ± 0	1097 ± 67	26 ± 1	39 ± 1
1.5 %	<3.8	102 ± 2	830 ± 17	16 ± 1	<21
2 %	<3.8	116 ± 1	973 ± 40	9 ± 1	<21
2 %	<3.8	131 ± 1	1111 ± 60	18 ± 1	26 ± 3
2.25 %	<3.8	107 ± 1	808 ± 20	20 ± 1	<21
2.25 %	<3.8	165 ± 1	1401 ± 42	19 ± 1	<21
2.5 %	<3.8	109 ± 2	917 ± 149	10 ± 0	<21
2.5 %	<3.8	129 ± 2	1283 ± 76	10 ± 1	32 ± 4
3 %	5.8 ± 0.7	119 ± 1	1028 ± 78	13 ± 0	30 ± 1
3 %	5.1 ± 0.8	124 ± 0	1220 ± 71	12 ± 1	<21
3.5 %	4.2 ± 0.8	125 ± 1	935 ± 71	10 ± 0	<21
3.5 %	<3.8	115 ± 1	890 ± 47	10 ± 1	<21
5 %	<3.8	236	1937	10	29 ± 0
5 %	<3.8	195	1568	<8	<21
10 %	<3.8	240 ± 1	2115 ± 19	9 ± 0	<21
10 %	<3.8	266 ± 1	2030 ± 56	18 ± 0	34 ± 0

**Table A-8a. Inductively Coupled Plasma-Mass Spectrometry Data from the Primary Adsorption Isotherm Experiment: NC Apatite (2 of 4)**

Weight % of NC-Apatite	Cesium µg/L	Strontium µg/L	Barium µg/L	Arsenic µg/L	Lead µg/L
0.5 %	<0.01	74.6 ± 0.5	13.0 ± 0.08	0.33 ± 0.05	0.39 ± 0.008
0.5 %	<0.01	61.2 ± 0.3	10.4 ± 0.05	0.32 ± 0.03	0.13 ± 0.012
1 %	<0.01	68.1 ± 0.5	11.3 ± 0.03	0.28 ± 0.04	0.25 ± 0.004
1 %	<0.01	70.6 ± 0.6	12.6 ± 0.13	0.33 ± 0.03	<0.1
1.5 %	<0.01	77.0 ± 0.2	12.5 ± 0.10	0.36 ± 0.03	<0.1
1.5 %	<0.01	66.7 ± 0.6	11.0 ± 0.03	0.36 ± 0.05	0.19 ± 0.007
2 %	<0.01	75.5 ± 0.4	11.6 ± 0.10	0.37 ± 0.03	<0.1
2 %	<0.01	76.5 ± 1.0	12.1 ± 0.14	0.41 ± 0.03	0.13 ± 0.013
2.25 %	0.026 ± 0.004	73.5 ± 0.6	11.3 ± 0.12	0.37 ± 0.04	0.31 ± 0.002
2.25 %	0.011 ± 0.001	88.2 ± 0.8	13.6 ± 0.03	0.42 ± 0.04	0.12 ± 0.004
2.5 %	<0.01	73.7 ± 0.6	11.0 ± 0.07	0.42 ± 0.05	<0.1
2.5 %	<0.01	82.2 ± 0.4	12.3 ± 0.04	0.41 ± 0.02	<0.1
3 %	<0.01	83.2 ± 0.9	11.8 ± 0.20	0.45 ± 0.06	<0.1
3 %	<0.01	81.6 ± 0.5	11.9 ± 0.12	0.41 ± 0.03	0.30 ± 0.008
3.5 %	<0.01	85.7 ± 1.0	12.9 ± 0.11	0.43 ± 0.04	<0.1
3.5 %	<0.01	80.4 ± 0.7	12.3 ± 0.05	0.40 ± 0.02	<0.1
5 %	<0.01	117	15.3	0.44	<0.1
5 %	<0.01	99.4	13.2	0.42	<0.1
10 %	<0.01	179 ± 1.9	15.5 ± 0.07	0.66 ± 0.00	<0.1
10 %	<0.01	174 ± 2.3	15.3 ± 0.11	0.54 ± 0.04	<0.1
NC Apatite	µmol/L	µmol/L	µmol/L	µmol/L	µmol/L
0.5 %	<0.08	851 ± 6	95 ± 1	4.4 ± 0.6	1.86 ± 0.04
0.5 %	<0.08	698 ± 3	75 ± 0	4.2 ± 0.3	0.65 ± 0.06
1 %	<0.08	777 ± 5	82 ± 0	3.8 ± 0.5	1.20 ± 0.02
1 %	<0.08	806 ± 6	92 ± 1	4.3 ± 0.3	<0.5
1.5 %	<0.08	878 ± 3	91 ± 1	4.8 ± 0.4	<0.5
1.5 %	<0.08	762 ± 7	80 ± 0	4.8 ± 0.7	0.92 ± 0.03
2 %	<0.08	862 ± 5	85 ± 1	4.9 ± 0.3	<0.5
2 %	<0.08	873 ± 12	88 ± 1	5.5 ± 0.3	0.62 ± 0.06
2.25 %	0.20 ± 0.03	839 ± 6	82 ± 1	5.0 ± 0.5	1.49 ± 0.01
2.25 %	0.08 ± 0.01	1007 ± 9	99 ± 0	5.6 ± 0.5	0.58 ± 0.02
2.5 %	<0.08	841 ± 7	80 ± 1	5.6 ± 0.7	<0.5
2.5 %	<0.08	938 ± 5	90 ± 0	5.5 ± 0.2	<0.5
3 %	<0.08	950 ± 10	86 ± 1	6.0 ± 0.8	<0.5
3 %	<0.08	931 ± 6	87 ± 1	5.5 ± 0.4	1.42 ± 0.04
3.5 %	<0.08	978 ± 12	94 ± 1	5.8 ± 0.6	<0.5
3.5 %	<0.08	918 ± 8	90 ± 0	5.3 ± 0.3	<0.5
5 %	<0.08	1339 ± 0	111 ± 0	5.9 ± 0.0	<0.5
5 %	<0.08	1134 ± 0	96 ± 0	5.6 ± 0.0	<0.5
10 %	<0.08	2048 ± 22	113 ± 1	8.8 ± 0.1	<0.5
10 %	<0.08	1991 ± 27	111 ± 1	7.2 ± 0.6	<0.5



**Table A-8a. Inductively Coupled Plasma-Mass Spectrometry Data from the Primary Adsorption Isotherm Experiment: NC Apatite (4 of 4)**

Weight % of NC Apatite	Gadolinium µg/L	Dysprosium µg/L	Holmium µg/L	Erbium µg/L	Ytterbium µg/L
0.5 %	<0.02	0.08 ± 0.01	0.03 ± 0.00	<0.02	<0.02
0.5 %	0.04 ± 0.00	<0.02	<0.02	<0.02	<0.02
1 %	0.03 ± 0.01	0.05 ± 0.01	0.02 ± 0.01	<0.02	<0.02
1 %	0.08 ± 0.02	0.04 ± 0.00	0.10 ± 0.01	0.04 ± 0.00	0.03 ± 0.01
1.5 %	<0.02	0.07 ± 0.02	<0.02	<0.02	<0.02
1.5 %	<0.02	0.11 ± 0.01	<0.02	<0.02	<0.02
2 %	0.04 ± 0.01	0.08 ± 0.01	<0.02	<0.02	<0.02
2 %	<0.02	0.14 ± 0.00	<0.02	<0.02	<0.02
2.25 %	<0.02	0.05 ± 0.01	0.03 ± 0.00	<0.02	<0.02
2.25 %	<0.02	0.04 ± 0.01	0.02 ± 0.00	<0.02	<0.02
2.5 %	<0.02	0.10 ± 0.02	0.02 ± 0.00	<0.02	<0.02
2.5 %	0.03 ± 0.01	0.04 ± 0.01	<0.02	0.03 ± 0.01	<0.02
3 %	<0.02	0.05 ± 0.00	<0.02	<0.02	<0.02
3 %	0.06 ± 0.01	0.26 ± 0.03	0.32 ± 0.01	<0.02	<0.02
3.5 %	<0.02	0.06 ± 0.00	0.03 ± 0.01	<0.02	<0.02
3.5 %	<0.02	<0.02	<0.02	<0.02	<0.02
5 %	<0.02	<0.02	<0.02	<0.02	<0.02
5 %	<0.02	0.04	<0.02	<0.02	<0.02
10 %	<0.02	<0.02	<0.02	<0.02	<0.02
10 %	<0.02	<0.02	<0.02	<0.02	<0.02
NC Apatite	µmol/L	µmol/L	µmol/L	µmol/L	µmol/L
0.5 %	<0.13	0.47 ± 0.07	0.20 ± 0.03	<0.12	<0.12
0.5 %	0.27 ± 0.01	<0.12	<0.12	<0.12	<0.12
1 %	0.21 ± 0.09	0.32 ± 0.07	0.15 ± 0.03	<0.12	<0.12
1 %	0.49 ± 0.12	0.27 ± 0.01	0.61 ± 0.06	0.24 ± 0.03	0.18 ± 0.05
1.5 %	<0.13	0.42 ± 0.12	<0.12	<0.12	<0.12
1.5 %	<0.13	0.70 ± 0.06	<0.12	<0.12	<0.12
2 %	0.23 ± 0.06	0.49 ± 0.05	<0.12	<0.12	<0.12
2 %	<0.13	0.84 ± 0.02	<0.12	<0.12	<0.12
2.25 %	<0.13	0.30 ± 0.04	0.21 ± 0.03	<0.12	<0.12
2.25 %	<0.13	0.23 ± 0.04	0.14 ± 0.02	<0.12	<0.12
2.5 %	<0.13	0.62 ± 0.14	0.13 ± 0.01	<0.12	<0.12
2.5 %	0.17 ± 0.09	0.23 ± 0.05	<0.12	0.16 ± 0.09	<0.12
3 %	<0.13	0.33 ± 0.03	<0.12	<0.12	<0.12
3 %	0.35 ± 0.08	1.62 ± 0.21	1.92 ± 0.03	<0.12	<0.12
3.5 %	<0.13	0.38 ± 0.02	0.16 ± 0.08	<0.12	<0.12
3.5 %	<0.13	<0.12	<0.12	<0.12	<0.12
5 %	<0.13	<0.12	<0.12	<0.12	<0.12
5 %	<0.13	0.25 ± 0.00	<0.12	<0.12	<0.12
10 %	<0.13	<0.12	<0.12	<0.12	<0.12
10 %	<0.13	<0.12	<0.12	<0.12	<0.12

**Table A-8b. Inductively Coupled Plasma-Mass Spectrometry Data from the Primary Adsorption Isotherm Experiment: Bone Char (1 of 4)**

Weight % of Bone Char	Chromium µg/L	Manganese µg/L	Iron µg/L	Copper µg/L	Zinc µg/L
0.5 %	0.46 ± 0.05	6.5 ± 0.10	78.1 ± 3.6	0.88 ± 0.02	<1.4
0.5 %	<0.2	7.1 ± 0.06	77.8 ± 6.0	<0.5	<1.4
1 %	<0.2	6.5 ± 0.03	62.9 ± 0.9	<0.5	<1.4
1 %	<0.2	9.5 ± 0.07	102 ± 5.1	0.84 ± 0.03	3.01 ± 0.05
1.5 %	<0.2	5.9 ± 0.02	62.2 ± 1.5	0.55 ± 0.04	1.45 ± 0.07
1.5 %	<0.2	3.5 ± 0.07	56.6 ± 2.9	0.51 ± 0.03	<1.4
2 %	<0.2	3.3 ± 0.02	28.6 ± 2.0	<0.5	1.40 ± 0.05
2 %	<0.2	2.5 ± 0.11	29.4 ± 3.5	0.97 ± 0.06	<1.4
2.25 %	<0.2	3.8 ± 0.14	47.7 ± 4.2	1.13 ± 0.05	2.23 ± 0.05
2.25 %	<0.2	3.6 ± 0.06	55.1 ± 2.2	0.57 ± 0.02	<1.4
2.5 %	0.21 ± 0.02	1.11 ± 0.04	na	1.14 ± 0.01	<1.4
2.5 %	<0.2	1.12 ± 0.02	na	0.65 ± 0.03	2.06 ± 0.07
3 %	<0.2	0.91 ± 0.04	na	0.56 ± 0.03	7.48 ± 0.12
3 %	<0.2	0.89 ± 0.00	na	0.54 ± 0.04	<1.4
3.5 %	0.27 ± 0.04	0.66 ± 0.03	na	1.94 ± 0.09	4.16 ± 0.06
3.5 %	0.33 ± 0.01	1.64 ± 0.02	na	<0.5	<1.4
5 %	0.64 ± 0.05	0.30 ± 0.01	na	0.60 ± 0.03	3.53 ± 0.17
5 %	0.56 ± 0.03	0.36 ± 0.02	na	0.90 ± 0.03	4.67 ± 0.11
10 %	1.31 ± 0.03	0.25 ± 0.01	na	<0.5	<1.4
10 %	1.38 ± 0.03	0.18 ± 0.01	na	<0.5	<1.4
Bone Char	µmol/L	µmol/L	µmol/L	µmol/L	µmol/L
0.5 %	8.9 ± 1.0	119 ± 1.8	1399 ± 64	13.9 ± 0.3	<21
0.5 %	<3.8	128 ± 1.1	1393 ± 107	<8	<21
1 %	<3.8	118 ± 0.6	1126 ± 16	<8	<21
1 %	<3.8	173 ± 1.2	1832 ± 91	13.2 ± 0.4	46 ± 1
1.5 %	<3.8	108 ± 0.3	1113 ± 28	8.6 ± 0.7	22 ± 1
1.5 %	<3.8	64.5 ± 1.3	1014 ± 52	8.0 ± 0.5	<21
2 %	<3.8	60.2 ± 0.3	513 ± 35	<8	21 ± 1
2 %	<3.8	45.3 ± 2.0	527 ± 62	15.3 ± 1.0	<21
2.25 %	<3.8	68.5 ± 2.6	854 ± 76	17.8 ± 0.7	34 ± 1
2.25 %	<3.8	65.5 ± 1.1	986 ± 39	8.9 ± 0.3	<21
2.5 %	4.0 ± 0.4	20.3 ± 0.7	na	17.9 ± 0.1	<21
2.5 %	<3.8	20.4 ± 0.5	na	10.3 ± 0.4	32 ± 1
3 %	<3.8	16.5 ± 0.7	na	8.9 ± 0.5	114 ± 2
3 %	<3.8	16.2 ± 0.0	na	8.5 ± 0.6	<21
3.5 %	5.3 ± 0.8	11.9 ± 0.5	na	30.5 ± 1.5	64 ± 1
3.5 %	6.3 ± 0.3	29.8 ± 0.4	na	<8 ±	<21
5 %	12.3 ± 1.1	5.5 ± 0.2	na	9.5 ± 0.5	54 ± 3
5 %	10.8 ± 0.6	6.6 ± 0.3	na	14.1 ± 0.4	71 ± 2
10 %	25.1 ± 0.6	4.6 ± 0.3	na	<8	<21
10 %	26.6 ± 0.6	3.3 ± 0.2	na	<8	<21

**Table A-8b. Inductively Coupled Plasma-Mass Spectrometry Data from the Primary Adsorption Isotherm Experiment: Bone Char (2 of 4)**

Weight % of Bone Char	Cesium μg/L	Strontium μg/L	Barium μg/L	Arsenic μg/L	Lead μg/L
0.5 %	<0.01	64.3 ± 0.4	12.2 ± 0.04	0.39 ± 0.05	0.11 ± 0.002
0.5 %	<0.01	62.2 ± 0.4	11.9 ± 0.05	0.41 ± 0.08	0.24 ± 0.002
1 %	<0.01	62.6 ± 0.1	12.5 ± 0.04	0.44 ± 0.01	<0.1
1 %	0.015 ± 0.003	70.8 ± 0.8	14.3 ± 0.04	0.40 ± 0.08	<0.1
1.5 %	<0.01	65.0 ± 0.8	13.8 ± 0.14	0.36 ± 0.03	<0.1
1.5 %	0.048 ± 0.004	54.3 ± 0.4	10.4 ± 0.10	0.38 ± 0.04	0.18 ± 0.01
2 %	<0.01	56.2 ± 0.2	11.3 ± 0.02	0.36 ± 0.02	<0.1
2 %	0.017 ± 0.003	50.0 ± 0.5	9.8 ± 0.13	0.29 ± 0.05	<0.1
2.25 %	0.013 ± 0.001	63.1 ± 0.4	13.5 ± 0.02	0.33 ± 0.04	<0.1
2.25 %	<0.01	58.0 ± 0.1	12.6 ± 0.02	0.35 ± 0.12	<0.1
2.5 %	<0.01	49.8 ± 0.4	10.0 ± 0.18	0.39 ± 0.02	<0.1
2.5 %	<0.01	50.9 ± 0.3	10.1 ± 0.06	0.36 ± 0.02	<0.1
3 %	<0.01	47.5 ± 0.8	9.7 ± 0.15	0.27 ± 0.02	0.23 ± 0.01
3 %	<0.01	49.4 ± 0.1	10.2 ± 0.11	0.32 ± 0.05	<0.1
3.5 %	0.015 ± 0.002	46.5 ± 0.7	9.9 ± 0.09	0.27 ± 0.03	<0.1
3.5 %	<0.01	50.0 ± 0.4	11.2 ± 0.06	0.30 ± 0.03	<0.1
5 %	<0.01	30.7 ± 0.2	10.7 ± 0.03	0.22 ± 0.01	<0.1
5 %	<0.01	36.1 ± 0.0	10.7 ± 0.09	0.22 ± 0.02	<0.1
10 %	<0.01	21.2 ± 0.2	10.1 ± 0.03	0.25 ± 0.03	<0.1
10 %	<0.01	21.6 ± 0.2	10.7 ± 0.11	0.29 ± 0.06	<0.1
Bone Char	μmol/L	μmol/L	μmol/L	μmol/L	μmol/L
0.5 %	<0.08	733 ± 4	88.8 ± 0.3	5.2 ± 0.6	0.55 ± 0.01
0.5 %	<0.08	710 ± 5	86.8 ± 0.4	5.5 ± 1.0	1.16 ± 0.01
1 %	<0.08	715 ± 1	90.9 ± 0.3	5.9 ± 0.1	<0.5
1 %	0.11 ± 0.02	808 ± 9	104 ± 0.3	5.4 ± 1.1	<0.5
1.5 %	<0.08	742 ± 9	100 ± 1.0	4.8 ± 0.4	<0.5
1.5 %	0.36 ± 0.03	620 ± 5	75.8 ± 0.7	5.1 ± 0.5	0.86 ± 0.03
2 %	<0.08	642 ± 2	82.0 ± 0.1	4.8 ± 0.3	<0.5
2 %	0.13 ± 0.02	570 ± 5	71.6 ± 0.9	3.9 ± 0.7	<0.5
2.25 %	0.10 ± 0.01	720 ± 5	98.1 ± 0.2	4.5 ± 0.5	<0.5
2.25 %	<0.08	662 ± 1	92.1 ± 0.1	4.6 ± 1.6	<0.5
2.5 %	<0.08	568 ± 4	72.5 ± 1.3	5.1 ± 0.3	<0.5
2.5 %	<0.08	581 ± 3	73.4 ± 0.5	4.8 ± 0.3	<0.5
3 %	<0.08	542 ± 9	70.9 ± 1.1	3.6 ± 0.2	1.13 ± 0.06
3 %	<0.08	564 ± 2	74.6 ± 0.8	4.2 ± 0.7	<0.5
3.5 %	0.11 ± 0.01	530 ± 8	72.1 ± 0.6	3.6 ± 0.4	<0.5
3.5 %	<0.08	570 ± 4	81.9 ± 0.4	4.1 ± 0.4	<0.5
5 %	<0.08	350 ± 2	77.7 ± 0.2	2.9 ± 0.1	<0.5
5 %	<0.08	412 ± 0	77.9 ± 0.7	2.9 ± 0.3	<0.5
10 %	<0.08	242 ± 2	73.4 ± 0.2	3.3 ± 0.5	<0.5
10 %	<0.08	247 ± 2	77.6 ± 0.8	3.8 ± 0.9	<0.5

**Table A-8b. Inductively Coupled Plasma-Mass Spectrometry Data from the Primary Adsorption Isotherm Experiment: Bone Char (3 of 4)**

Weight % of Bone Char	Yttrium µg/L	Lanthanum µg/L	Praeseodymium µg/L	Samarium µg/L	Europium µg/L
0.5 %	0.67 ± 0.01	0.05 ± 0.02	<0.05	<0.05	<0.05
0.5 %	0.16 ± 0.02	0.13 ± 0.04	<0.05	<0.05	<0.05
1 %	0.13 ± 0.03	0.18 ± 0.03	<0.05	<0.05	0.06 ± 0.01
1 %	0.20 ± 0.03	0.14 ± 0.03	<0.05	<0.05	<0.05
1.5 %	0.09 ± 0.03	0.08 ± 0.02	<0.05	<0.05	0.07 ± 0.01
1.5 %	0.20 ± 0.01	0.22 ± 0.03	0.05 ± 0.01	<0.05	0.07 ± 0.01
2 %	0.59 ± 0.02	0.06 ± 0.02	<0.05	<0.05	<0.05
2 %	0.12 ± 0.01	0.08 ± 0.01	<0.05	<0.05	<0.05
2.25 %	0.08 ± 0.01	0.06 ± 0.00	<0.05	<0.05	<0.05
2.25 %	0.25 ± 0.03	0.21 ± 0.03	0.06 ± 0.01	<0.05	0.07 ± 0.00
2.5 %	<0.06	0.08 ± 0.02	<0.05	<0.05	0.07 ± 0.02
2.5 %	<0.06	<0.05	<0.05	<0.05	0.09 ± 0.01
3 %	<0.06	<0.05	<0.05	<0.05	<0.05
3 %	<0.06	<0.05	<0.05	<0.05	0.08 ± 0.02
3.5 %	<0.06	<0.05	<0.05	<0.05	0.15 ± 0.01
3.5 %	<0.06	<0.05	<0.05	<0.05	<0.05
5 %	<0.06	<0.05	<0.05	<0.05	0.20 ± 0.01
5 %	<0.06	<0.05	<0.05	<0.05	<0.05
10 %	<0.06	<0.05	<0.05	<0.05	<0.05
10 %	0.26 ± 0.01	0.23 ± 0.01	0.05 ± 0.01	<0.05	0.10 ± 0.02
Bone Char	µmol/L	µmol/L	µmol/L	µmol/L	µmol/L
0.5 %	7.6 ± 0.1	0.4 ± 0.1	<0.35	<0.3	<0.33
0.5 %	1.8 ± 0.2	0.9 ± 0.3	<0.35	<0.3	<0.33
1 %	1.4 ± 0.3	1.3 ± 0.2	<0.35	<0.3	0.39 ± 0.08
1 %	2.3 ± 0.4	1.0 ± 0.2	<0.35	<0.3	<0.33
1.5 %	1.0 ± 0.3	0.6 ± 0.1	<0.35	<0.3	0.49 ± 0.04
1.5 %	2.2 ± 0.1	1.6 ± 0.2	0.39 ± 0.07	<0.3	0.43 ± 0.05
2 %	6.6 ± 0.3	0.4 ± 0.1	<0.35	<0.3	<0.33
2 %	1.4 ± 0.1	0.6 ± 0.1	<0.35	<0.3	<0.33
2.25 %	0.9 ± 0.1	0.4 ± 0.0	<0.35	<0.3	<0.33
2.25 %	2.9 ± 0.4	1.5 ± 0.2	0.43 ± 0.07	<0.3	0.45 ± 0.01
2.5 %	<0.7	0.6 ± 0.2	<0.35	<0.3	0.43 ± 0.13
2.5 %	<0.7	<0.4	<0.35	<0.3	0.59 ± 0.05
3 %	<0.7	<0.4	<0.35	<0.3	<0.33
3 %	<0.7	<0.4	<0.35	<0.3	0.53 ± 0.11
3.5 %	<0.7	<0.4	<0.35	<0.3	0.97 ± 0.04
3.5 %	<0.7	<0.4	<0.35	<0.3	<0.33
5 %	<0.7	<0.4	<0.35	<0.3	1.30 ± 0.06
5 %	<0.7	<0.4	<0.35	<0.3	<0.33
10 %	<0.7	<0.4	<0.35	<0.3	<0.33
10 %	2.9 ± 0.1	1.7 ± 0.1	0.39 ± 0.05	<0.3	0.69 ± 0.11

**Table A-8b. Inductively Coupled Plasma-Mass Spectrometry Data from the Primary Adsorption Isotherm Experiment: Bone Char (4 of 4)**

Weight % of Bone Char	Gadolinium		Dysprosium		Holmium		Erbium		Ytterbium	
	µg/L		µg/L		µg/L		µg/L		µg/L	
0.5 %	0.09	± 0.005	7.40	± 0.06	0.06	± 0.003	0.07	± 0.01	<0.02	
0.5 %	<0.02		<0.02		<0.02		<0.02		<0.02	
1 %	<0.02		0.09	± 0.004	0.12	± 0.002	<0.02		<0.02	
1 %	<0.02		<0.02		<0.02		0.03	± 0.01	<0.02	
1.5 %	<0.02		<0.02		<0.02		<0.02		<0.02	
1.5 %	<0.02		<0.02		0.03	± 0.01	0.02	± 0.01	<0.02	
2 %	0.06	± 0.01	7.75	± 0.08	0.05	± 0.01	0.03	± 0.01	<0.02	
2 %	<0.02		0.03	± 0.00	<0.02		<0.02		<0.02	
2.25 %	<0.02		0.09	± 0.01	<0.02		<0.02		<0.02	
2.25 %	0.03	± 0.02	0.06	± 0.01	0.02	± 0.01	0.04	± 0.01	<0.02	
2.5 %	<0.02		0.04	± 0.01	0.03	± 0.01	<0.02		<0.02	
2.5 %	<0.02		<0.02		<0.02		<0.02		<0.02	
3 %	<0.02		<0.02		<0.02		<0.02		<0.02	
3 %	<0.02		0.06	± 0.004	0.03	± 0.01	<0.02		<0.02	
3.5 %	<0.02		0.02	± 0.002	<0.02		<0.02		<0.02	
3.5 %	<0.02		0.04	± 0.01	<0.02		<0.02		<0.02	
5 %	<0.02		<0.02		<0.02		<0.02		<0.02	
5 %	0.03	± 0.02	0.03	± 0.01	<0.02		<0.02		<0.02	
10 %	0.05	± 0.01	0.48	± 0.02	<0.02		<0.02		<0.02	
10 %	0.09	± 0.03	0.57	± 0.02	0.27	± 0.01	<0.02		<0.02	
Bone Char	µmol/L		µmol/L		µmol/L		µmol/L		µmol/L	
0.5 %	0.56	± 0.03	45.56	± 0.37	0.38	± 0.02	0.39	± 0.05	<0.12	
0.5 %	<0.13		<0.12		<0.12		<0.12		<0.12	
1 %	<0.13		0.53	± 0.03	0.70	± 0.01	<0.12		<0.12	
1 %	<0.13		<0.12		<0.12		0.16	± 0.05	<0.12	
1.5 %	<0.13		<0.12		<0.12		<0.12		<0.12	
1.5 %	<0.13		<0.12		0.16	± 0.06	0.14	± 0.08	<0.12	
2 %	0.40	± 0.09	47.68	± 0.51	0.32	± 0.05	0.17	± 0.08	<0.12	
2 %	<0.13		0.22	± 0.02	<0.12		<0.12		<0.12	
2.25 %	<0.13		0.54	± 0.04	<0.12		<0.12		<0.12	
2.25 %	0.17	± 0.14	0.37	± 0.03	0.15	± 0.06	0.21	± 0.07	<0.12	
2.5 %	<0.13		0.23	± 0.04	0.21	± 0.03	<0.12		<0.12	
2.5 %	<0.13		<0.12		<0.12		<0.12		<0.12	
3 %	<0.13		<0.12		<0.12		<0.12		<0.12	
3 %	<0.13		0.34	± 0.02	0.18	± 0.04	<0.12		<0.12	
3.5 %	<0.13		0.13	± 0.01	<0.12		<0.12		<0.12	
3.5 %	<0.13		0.23	± 0.06	<0.12		<0.12		<0.12	
5 %	<0.13		<0.12		<0.12		<0.12		<0.12	
5 %	0.21	± 0.11	0.19	± 0.08	<0.12		<0.12		<0.12	
10 %	0.30	± 0.06	2.96	± 0.13	<0.12		<0.12		<0.12	
10 %	0.57	± 0.16	3.52	± 0.11	1.64	± 0.06	<0.12		<0.12	







**Table A-8c. Inductively Coupled Plasma-Mass Spectrometry Data from the Primary Adsorption Isotherm Experiment: Hydroxyapatite (4 of 4)**

Weight % of Hydroxyapatite	Gadolinium µg/L	Dysprosium µg/L	Holmium µg/L	Erbium µg/L	Ytterbium µg/L
0.5 %	<0.02	<0.02	<0.02	<0.02	<0.02
0.5 %	0.09 ± 0.01	0.24 ± 0.01	<0.02	<0.02	<0.02
1 %	<0.02	0.13 ± 0.01	<0.02	<0.02	<0.02
1 %	<0.02	<0.02	<0.02	<0.02	<0.02
1.5 %	<0.02	<0.02	0.03 ± 0.01	<0.02	<0.02
1.5 %	<0.02	<0.02	<0.02	<0.02	<0.02
2 %	<0.02	<0.02	<0.02	<0.02	<0.02
2 %	<0.02	<0.02	<0.02	<0.02	<0.02
2.25 %	<0.02	<0.02	<0.02	<0.02	<0.02
2.25 %	<0.02	0.04 ± 0.02	<0.02	<0.02	<0.02
2.5 %	<0.02	0.04 ± 0.01	0.05 ± 0.01	<0.02	<0.02
2.5 %	<0.02	<0.02	<0.02	<0.02	<0.02
3 %	<0.02	<0.02	<0.02	<0.02	<0.02
3 %	<0.02	<0.02	<0.02	<0.02	<0.02
3.5 %	<0.02	0.02 ± 0.01	<0.02	<0.02	<0.02
3.5 %	<0.02	0.05 ± 0.01	<0.02	<0.02	<0.02
5 %	<0.02	<0.02	<0.02	<0.02	<0.02
5 %	0.08 ± 0.01	0.04 ± 0.02	0.02 ± 0.00	<0.02	<0.02
10 %	<0.02	<0.02	<0.02	<0.02	<0.02
10 %	<0.02	<0.02	<0.02	<0.02	<0.02
Hydroxyapatite	µmol/L	µmol/L	µmol/L	µmol/L	µmol/L
0.5 %	<0.13	<0.12	<0.12	<0.12	<0.12
0.5 %	0.57 ± 0.06	1.45 ± 0.05	<0.12	<0.12	<0.12
1 %	<0.13	0.77 ± 0.06	<0.12	<0.12	<0.12
1 %	<0.13	<0.12	<0.12	<0.12	<0.12
1.5 %	<0.13	<0.12	0.19 ± 0.03	<0.12	<0.12
1.5 %	<0.13	<0.12	<0.12	<0.12	<0.12
2 %	<0.13	<0.12	<0.12	<0.12	<0.12
2 %	<0.13	<0.12	<0.12	<0.12	<0.12
2.25 %	<0.13	<0.12	<0.12	<0.12	<0.12
2.25 %	<0.13	0.23 ± 0.09	<0.12	<0.12	<0.12
2.5 %	<0.13	0.22 ± 0.05	0.30 ± 0.04	<0.12	<0.12
2.5 %	<0.13	<0.12	<0.12	<0.12	<0.12
3 %	<0.13	<0.12	<0.12	<0.12	<0.12
3 %	<0.13	<0.12	<0.12	<0.12	<0.12
3.5 %	<0.13	0.15 ± 0.07	<0.12	<0.12	<0.12
3.5 %	<0.13	0.33 ± 0.04	<0.12	<0.12	<0.12
5 %	<0.13	<0.12	<0.12	<0.12	<0.12
5 %	0.53 ± 0.06	0.25 ± 0.10	0.13 ± 0.03	<0.12	<0.12
10 %	<0.13	<0.12	<0.12	<0.12	<0.12
10 %	<0.13	<0.12	<0.12	<0.12	<0.12

**Table A-8d. Inductively Coupled Plasma-Mass Spectrometry Data from the Primary Adsorption Isotherm Experiment: Clinoptilolite (1 of 4)**

Weight % of Clinoptilolite	Chromium µg/L	Manganese µg/L	Iron µg/L	Copper µg/L	Zinc µg/L
0.5 %	<0.2	3.9 ± 0.1	31.5 ± 3.3	1.68 ± 0.15	<1.4
0.5 %	<0.2	5.1 ± 0.1	58.2 ± 0.7	7.79 ± 0.02	1.9 ± 0.1
1 %	<0.2	4.4 ± 0.0	59.4 ± 2.2	0.93 ± 0.07	<1.4
1 %	<0.2	5.5 ± 0.1	78.3 ± 4.9	<0.50	<1.4
1.5 %	<0.2	2.5 ± 0.0	13.2 ± 1.4	<0.50	<1.4
1.5 %	<0.2	2.8 ± 0.0	21.8 ± 3.8	0.81 ± 0.05	12.2 ± 0.3
2 %	0.25 ± 0.04	6.7 ± 0.1	118 ± 3.8	<0.50	3.1 ± 0.3
2 %	<0.2	4.2 ± 0.1	58.3 ± 4.1	<0.50	<1.4
2.25 %	<0.2	2.7 ± 0.0	27.1 ± 3.5	<0.50	<1.4
2.25 %	<0.2	8.7 ± 0.1	175 ± 5.1	0.57 ± 0.04	<1.4
2.5 %	0.75 ± 0.04	40.6 ± 0.2	1089 ± 34	0.71 ± 0.02	1.5 ± 0.03
2.5 %	<0.2	2.4 ± 0.0	31.5 ± 3.0	<0.50	2.4 ± 0.3
3 %	<0.2	2.9 ± 0.0	48.0 ± 3.7	0.90 ± 0.07	<1.4
3 %	<0.2	5.6 ± 0.0	102 ± 4.0	14.26 ± 0.18	<1.4
3.5 %	<0.2	3.80 ± 0.0	67.9 ± 1.2	<0.50	<1.4
3.5 %	1.14 ± 0.03	7.75 ± 0.1	206 ± 3.3	1.31 ± 0.03	<1.4
5 %	<0.2	1.24 ± 0.0	13.7 ± 3.3	<0.50	<1.4
5 %	<0.2	1.76 ± 0.0	19.6 ± 1.1	<0.50	<1.4
10 %	<2.0	16.8 ± 0.3	376 ± 38	6.17 ± 0.31	127 ± 2
10 %	<2.0	19.7 ± 0.2	497 ± 23	<0.50	<1.4
Clinoptilolite	µmol/L	µmol/L	µmol/L	µmol/L	µmol/L
0.5 %	< 3.8	70.4 ± 1.5	564 ± 59	26.5 ± 2.4	<21
0.5 %	< 3.8	92.6 ± 2.2	1041 ± 13	122.6 ± 0.3	29 ± 2
1 %	< 3.8	80.4 ± 0.8	1063 ± 40	14.7 ± 1.1	<21
1 %	< 3.8	99.4 ± 2.1	1401 ± 88	< 8.0	<21
1.5 %	< 3.8	45.6 ± 0.3	236 ± 26	< 8.0	<21
1.5 %	< 3.8	51.4 ± 0.7	391 ± 69	12.7 ± 0.7	186 ± 4
2 %	4.8 ± 0.7	121.5 ± 1.5	2117 ± 69	< 8.0	48 ± 5
2 %	< 3.8	77.0 ± 1.5	1044 ± 74	< 8.0	<21
2.25 %	< 3.8	48.8 ± 0.7	486 ± 62	< 8.0	<21
2.25 %	< 3.8	158.5 ± 1.1	3133 ± 92	9.0 ± 0.6	<21
2.5 %	14.4 ± 0.7	739.1 ± 4.0	19508 ± 609	11.1 ± 0.3	23 ± 0
2.5 %	< 3.8	43.1 ± 0.3	564 ± 54	< 8.0	37 ± 4
3 %	< 3.8	52.1 ± 0.8	860 ± 66	14.2 ± 1.1	<21
3 %	< 3.8	101.7 ± 0.7	1831 ± 72	224.3 ± 2.8	<21
3.5 %	< 3.8	69.2 ± 0.5	1216 ± 21	< 8.0	<21
3.5 %	22.0 ± 0.5	141.1 ± 1.6	3680 ± 59	20.6 ± 0.5	<21
5 %	< 3.8	22.5 ± 0.9	246 ± 60	< 8.0	<21
5 %	< 3.8	32.1 ± 0.8	350 ± 19	< 8.0	<21
10 %	< 3.8	305.6 ± 6.0	6728 ± 675	97.0 ± 4.9	1950 ± 32
10 %	< 3.8	359.4 ± 4.4	8896 ± 409	< 8.0	<21

**Table A-8d. Inductively Coupled Plasma-Mass Spectrometry Data from the Primary Adsorption Isotherm Experiment: Clinoptilolite (2 of 4)**

Weight % of Clinoptilolite	Cesium µg/L	Strontium µg/L	Barium µg/L	Arsenic µg/L	Lead µg/L
0.5 %	<0.01	30.9 ± 0.4	7.5 ± 0.1	0.76 ± 0.05	0.25 ± 0.01
0.5 %	0.15 ± 0.002	32 ± 0.3	8.0 ± 0.1	0.62 ± 0.04	<0.1
1 %	<0.01	20.1 ± 0.1	5.5 ± 0.1	0.84 ± 0.04	0.23 ± 0.004
1 %	0.64 ± 0.02	21 ± 0.3	6.2 ± 0.1	0.86 ± 0.07	<0.1
1.5 %	0.12 ± 0.01	20.4 ± 0.1	4.9 ± 0.1	0.78 ± 0.10	<0.1
1.5 %	0.09 ± 0.003	17.1 ± 0.1	4.6 ± 0.01	0.97 ± 0.14	0.25 ± 0.01
2 %	<0.01	17.5 ± 0.1	5.5 ± 0.05	1.06 ± 0.04	<0.1
2 %	<0.01	16.1 ± 0.2	4.3 ± 0.1	0.94 ± 0.08	<0.1
2.25 %	<0.01	13.0 ± 0.1	3.2 ± 0.02	0.77 ± 0.05	0.52 ± 0.005
2.25 %	<0.01	15.2 ± 0.1	4.9 ± 0.04	0.92 ± 0.05	0.35 ± 0.02
2.5 %	<0.01	9.7 ± 0.3	10.7 ± 0.1	1.04 ± 0.10	0.42 ± 0.01
2.5 %	<0.01	9.1 ± 0.1	2.4 ± 0.04	0.85 ± 0.02	<0.1
3 %	<0.01	8.6 ± 0.1	2.3 ± 0.03	1.00 ± 0.06	0.16 ± 0.01
3 %	<0.01	9.4 ± 0.1	4.1 ± 0.05	0.86 ± 0.08	<0.1
3.5 %	0.03 ± 0.02	7.3 ± 0.1	2.5 ± 0.02	1.02 ± 0.05	<0.1
3.5 %	0.06 ± 0.01	7.5 ± 0.03	3.5 ± 0.03	1.07 ± 0.05	<0.1
5 %	<0.01	0.4 ± 0.02	0.4 ± 0.00	0.16 ± 0.04	<0.1
5 %	<0.01	0.4 ± 0.01	0.4 ± 0.01	0.11 ± 0.02	<0.1
10 %	na	5 ± 0.2	7.9 ± 0.2	3.66 ± 0.99	5.88 ± 0.13
10 %	<0.01	3.5 ± 0.2	5.5 ± 0.1	4.15 ± 0.29	<0.1
Clinoptilolite	µmol/L	µmol/L	µmol/L	µmol/L	µmol/L
0.5 %	<0.08	353 ± 4	54.5 ± 0.8	10.1 ± 0.7	1.2 ± 0.1
0.5 %	1.10 ± 0.02	360 ± 4	57.9 ± 0.5	8.2 ± 0.5	<0.5
1 %	<0.08	229 ± 1	40.2 ± 0.4	11.2 ± 0.6	1.1 ± 0.02
1 %	4.85 ± 0.15	237 ± 3	45.2 ± 0.5	11.5 ± 1.0	<0.5
1.5 %	0.91 ± 0.06	233 ± 1	35.8 ± 0.4	10.4 ± 1.4	<0.5
1.5 %	0.69 ± 0.03	195 ± 1	33.9 ± 0.1	13.0 ± 1.8	1.2 ± 0.04
2 %	<0.08	199 ± 1	40.2 ± 0.3	14.2 ± 0.5	<0.5
2 %	<0.08	184 ± 3	31.5 ± 0.9	12.6 ± 1.1	<0.5
2.25 %	<0.08	148 ± 1	23.3 ± 0.1	10.3 ± 0.6	2.5 ± 0.02
2.25 %	<0.08	174 ± 1	36.0 ± 0.3	12.3 ± 0.7	1.7 ± 0.1
2.5 %	<0.08	110 ± 3	77.8 ± 0.7	13.8 ± 1.3	2.0 ± 0.04
2.5 %	<0.08	104 ± 1	17.5 ± 0.3	11.3 ± 0.3	<0.5
3 %	<0.08	98 ± 1	16.8 ± 0.2	13.3 ± 0.7	0.8 ± 0.1
3 %	<0.08	107 ± 1	30.0 ± 0.4	11.5 ± 1.1	<0.5
3.5 %	0.22 ± 0.19	83 ± 1	18.2 ± 0.2	13.6 ± 0.7	<0.5
3.5 %	0.44 ± 0.07	86 ± 0	25.6 ± 0.2	14.3 ± 0.7	<0.5
5 %	<0.08	5.1 ± 0.2	2.9 ± 0.0	2.2 ± 0.5	<0.5
5 %	<0.08	5.1 ± 0.2	3.1 ± 0.1	1.5 ± 0.3	<0.5
10 %	0.00 ± 0.00	55 ± 2	57.2 ± 1.4	49 ± 13	28.4 ± 0.6
10 %	<0.08	40 ± 3	39.8 ± 0.8	55 ± 4	<0.5



**Table A-8d. Inductively Coupled Plasma-Mass Spectrometry Data from the Primary Adsorption Isotherm Experiment: Clinoptilolite (4 of 4)**

Weight % of Clinoptilolite	Gadolinium μg/L	Dysprosium μg/L	Holmium μg/L	Erbium μg/L	Ytterbium μg/L
0.5 %	<0.02	0.07 ± 0.01	0.02 ± 0.002	<0.02	<0.02
0.5 %	0.11 ± 0.03	0.06 ± 0.004	0.02 ± 0.01	0.03 ± 0.003	<0.02
1 %	0.12 ± 0.04	0.14 ± 0.01	0.05 ± 0.001	0.05 ± 0.02	0.03 ± 0.01
1 %	0.13 ± 0.004	0.17 ± 0.01	0.04 ± 0.01	0.06 ± 0.02	0.05 ± 0.02
1.5 %	0.03 ± 0.03	0.06 ± 0.02	<0.02	<0.02	<0.02
1.5 %	0.06 ± 0.04	0.06 ± 0.04	<0.02	0.03 ± 0.01	<0.02
2 %	0.11 ± 0.06	0.09 ± 0.01	0.03 ± 0.01	0.06 ± 0.02	0.07 ± 0.02
2 %	0.06 ± 0.03	0.06 ± 0.01	0.02 ± 0.01	0.03 ± 0.002	<0.02
2.25 %	0.05 ± 0.01	0.08 ± 0.01	<0.02	<0.02	<0.02
2.25 %	0.19 ± 0.05	0.21 ± 0.02	0.06 ± 0.01	0.09 ± 0.02	0.13 ± 0.03
2.5 %	1.03 ± 0.12	0.63 ± 0.02	0.17 ± 0.005	0.48 ± 0.02	0.46 ± 0.11
2.5 %	<0.02	0.04 ± 0.01	<0.02	<0.02	<0.02
3 %	0.08 ± 0.03	0.11 ± 0.02	0.05 ± 0.003	0.04 ± 0.01	0.04 ± 0.04
3 %	0.09 ± 0.005	0.06 ± 0.03	0.02 ± 0.01	0.07 ± 0.02	0.04 ± 0.04
3.5 %	0.05 ± 0.03	0.09 ± 0.01	0.03 ± 0.000	0.05 ± 0.01	<0.02
3.5 %	0.14 ± 0.01	0.09 ± 0.01	0.05 ± 0.01	0.12 ± 0.02	0.06 ± 0.04
5 %	0.04 ± 0.01	0.12 ± 0.005	0.09 ± 0.01	0.03 ± 0.02	<0.02
5 %	<0.02	0.04 ± 0.01	<0.02	<0.02	<0.02
10 %	0.69 ± 0.46	0.87 ± 0.23	<0.2	0.26 ± 0.08	0.20 ± 0.03
10 %	0.60 ± 0.28	11.17 ± 0.66	<0.2	0.30 ± 0.19	<0.2
Clinoptilolite	μmol/L	μmol/L	μmol/L	μmol/L	μmol/L
0.5 %	<0.1	0.43 ± 0.06	0.13 ± 0.01	<0.12	<0.12
0.5 %	0.7 ± 0.2	0.36 ± 0.03	0.12 ± 0.04	0.2 ± 0.0	<0.12
1 %	0.8 ± 0.2	0.84 ± 0.06	0.29 ± 0.01	0.3 ± 0.1	0.15 ± 0.06
1 %	0.8 ± 0.0	1.04 ± 0.07	0.25 ± 0.04	0.3 ± 0.1	0.28 ± 0.09
1.5 %	0.2 ± 0.2	0.34 ± 0.13	<0.12	<0.12	<0.12
1.5 %	0.4 ± 0.3	0.39 ± 0.22	<0.12	0.2 ± 0.0	<0.12
2 %	0.7 ± 0.4	0.56 ± 0.07	0.16 ± 0.06	0.3 ± 0.1	0.39 ± 0.13
2 %	0.4 ± 0.2	0.35 ± 0.09	0.13 ± 0.05	0.2 ± 0.0	<0.12
2.25 %	0.3 ± 0.1	0.47 ± 0.07	<0.12	<0.12	<0.12
2.25 %	1.2 ± 0.3	1.29 ± 0.12	0.35 ± 0.04	0.5 ± 0.1	0.77 ± 0.18
2.5 %	6.5 ± 0.7	3.91 ± 0.14	1.02 ± 0.03	2.9 ± 0.1	2.66 ± 0.61
2.5 %	<0.1	0.26 ± 0.06	<0.12	<0.12	<0.12
3 %	0.5 ± 0.2	0.67 ± 0.10	0.28 ± 0.02	0.2 ± 0.1	0.26 ± 0.24
3 %	0.6 ± 0.0	0.35 ± 0.18	0.14 ± 0.03	0.4 ± 0.1	0.25 ± 0.23
3.5 %	0.3 ± 0.2	0.53 ± 0.08	0.16 ± 0.00	0.3 ± 0.0	<0.12
3.5 %	0.9 ± 0.1	0.54 ± 0.08	0.28 ± 0.07	0.7 ± 0.1	0.38 ± 0.25
5 %	0.3 ± 0.1	0.71 ± 0.03	0.56 ± 0.09	0.2 ± 0.1	<0.12
5 %	<0.1	0.27 ± 0.03	<0.12	<0.12	<0.12
10 %	4.4 ± 2.9	5.35 ± 1.45	<0.12	1.5 ± 0.5	1.14 ± 0.16
10 %	3.8 ± 1.8	68.74 ± 4.07	<0.12	1.8 ± 1.1	<0.12

**Table A-8e. Inductively Coupled Plasma-Mass Spectrometry Data from the Primary Adsorption Isotherm Experiment: Hanford Soil (1 of 4)**

Weight % of Hanford Soil	Chromium µg/L	Manganese µg/L	Iron µg/L		Copper µg/L		Zinc µg/L
100 %	<0.2	4.9 ± 0.13	80.0	± 3.1	0.73	± 0.02	<1.4
100 %	<0.2	4.7 ± 0.10	68.2	± 2.6	<0.5		<1.4
100 %	<0.2	5.5 ± 0.05	30.0	± 4.8	0.79	± 0.05	1.69 ± 0.06
100 %	<0.2	5.3 ± 0.06	34.3	± 4.3	0.82	± 0.04	4.14 ± 0.08
100 %	<0.2	0.17 ± 0.01	na		0.83	± 0.02	3.36 ± 0.11
100 %	<0.2	0.18 ± 0.04	75.1	± 18.4	2.68	± 0.78	6.65 ± 1.37
100 %	<0.2	0.22 ± 0.01	86.4	± 1.8	2.31	± 0.04	2.22 ± 0.08
100 %	0.31 ± 0.02	0.19 ± 0.01	57.2	± 2.3	1.99	± 0.04	1.93 ± 0.04
100 %	<0.2	0.19 ± 0.02	na		2.00	± 0.04	3.13 ± 0.01
Avg. ± Std. Dev.	0.31 ± 0.02	2.38 ± 2.61	61.6	± 22.2	1.52	± 0.80	3.30 ± 1.71
Hanford Soil	µmol/L	µmol/L	µmol/L		µmol/L		µmol/L
100 %	<3.8	89.8 ± 2.4	1433	± 56	11.5	± 0.4	<21
100 %	<3.8	85.6 ± 1.8	1222	± 46	<8.0		<21
100 %	<3.8	100.3 ± 1.0	537	± 85	12.5	± 0.8	26 ± 1
100 %	<3.8	97.1 ± 1.0	614	± 77	12.9	± 0.6	63 ± 1
100 %	<3.8	3.0 ± 0.1	na		13.1	± 0.4	51 ± 2
100 %	<3.8	3.2 ± 0.7	1345	± 330	42.2	± 12.3	102 ± 21
100 %	<3.8	4.0 ± 0.1	1548	± 32	36.4	± 0.6	34 ± 1
100 %	6.0 ± 0.3	3.4 ± 0.2	1024	± 41	31.3	± 0.6	29 ± 1
100 %	<3.8	3.5 ± 0.3	na		31.5	± 0.7	48 ± 0

**Table A-8e. Inductively Coupled Plasma-Mass Spectrometry Data from the Primary Adsorption Isotherm Experiment: Hanford Soil (2 of 4)**

Weight % of Hanford Soil	Cesium µg/L	Strontium		Barium		Arsenic		Lead	
		µg/L	µg/L	µg/L	µg/L	µg/L	µg/L	µg/L	µg/L
100 %	<0.01	55.0	± 0.4	10.8	± 0.08	0.41	± 0.03	<0.1	
100 %	<0.01	56.8	± 0.1	10.5	± 0.12	0.37	± 0.01	<0.1	
100 %	<0.01	57.7	± 0.1	11.2	± 0.13	0.47	± 0.03	<0.1	
100 %	<0.01	61.9	± 0.3	11.9	± 0.01	0.46	± 0.09	0.20	± 0.01
100 %	<0.01	5.0	± 0.1	0.25	± 0.01	<0.03		0.17	± 0.01
100 %	<0.01	6.3	± 1.1	0.54	± 0.08	<0.03		1.54	± 0.32
100 %	<0.01	5.3	± 0.01	0.26	± 0.01	<0.03		0.35	± 0.01
100 %	<0.01	5.4	± 0.04	0.28	± 0.02	<0.03		0.25	± 0.02
100 %	<0.01	5.5	± 0.03	0.30	± 0.01	<0.03		0.51	± 0.01
Avg. ± Std. Dev.	0.01	28.76	± 27.65	5.11	± 5.69	0.43	± 0.05	0.50	± 0.52
Hanford Soil	µmol/L	µmol/L	µmol/L	µmol/L	µmol/L	µmol/L	µmol/L	µmol/L	µmol/L
100 %	<0.08	628	± 4	78.7	± 0.6	5.5	± 0.3	<0.5	
100 %	<0.08	648	± 1	76.3	± 0.8	4.9	± 0.2	<0.5	
100 %	<0.08	659	± 1	81.7	± 0.9	6.3	± 0.4	<0.5	
100 %	<0.08	706	± 4	86.5	± 0.1	6.1	± 1.2	1.0	± 0.0
100 %	<0.08	57	± 1	1.8	± 0.1	<0.4		0.8	± 0.0
100 %	<0.08	72	± 13	4.0	± 0.5	<0.4		7.4	± 1.6
100 %	<0.08	61	± 0.12	1.9	± 0.0	<0.4		1.7	± 0.0
100 %	<0.08	61	± 0.47	2.0	± 0.1	<0.4		1.2	± 0.1
100 %	<0.08	62	± 0.30	2.2	± 0.1	<0.4		2.5	± 0.1

**Table A-8e. Inductively Coupled Plasma-Mass Spectrometry Data from the Primary Adsorption Isotherm Experiment: Hanford Soil (3 of 4)**

Weight % of Hanford Soil	Yttrium µg/L	Lanthanum µg/L	Praeseodymium µg/L	Samarium µg/L	Europium µg/L
100 %	<0.06	<0.05	<0.05	<0.05	<0.05
100 %	<0.06	<0.05	<0.05	<0.05	<0.05
100 %	<0.06	<0.05	<0.05	<0.05	<0.05
100 %	0.08 ± 0.01	<0.05	<0.05	<0.05	<0.05
100 %	0.09 ± 0.02	<0.05	<0.05	<0.05	<0.05
100 %	0.06 ± 0.01	0.06 ± 0.04	<0.05	0.09 ± 0.04	<0.05
100 %	0.06	0.18 ± 0.02	<0.05	<0.05	<0.05
100 %	<0.06	0.06 ± 0.01	<0.05	<0.05	<0.05
100 %	0.12 ± 0.01	0.08 ± 0.01	<0.05	<0.05	<0.05

Avg. ± Std. Dev. 0.08 ± 0.03 0.10 ± 0.06

Hanford Soil	µmol/L	µmol/L	µmol/L	µmol/L	µmol/L
100 %	< 0.7	< 0.4	< 0.4	< 0.3	< 0.33
100 %	< 0.7	< 0.4	< 0.4	< 0.3	< 0.33
100 %	< 0.7	< 0.4	< 0.4	< 0.3	< 0.33
100 %	0.9 ± 0.1	< 0.4	< 0.4	< 0.3	< 0.33
100 %	1.1 ± 0.2	< 0.4	< 0.4	< 0.3	< 0.33
100 %	0.7 ± 0.1	0.4 ± 0.3	< 0.4	0.6 ± 0.2	< 0.33
100 %	< 0.7	1.3 ± 0.1	< 0.4	< 0.3	< 0.33
100 %	< 0.7	0.4 ± 0.1	< 0.4	< 0.3	< 0.33
100 %	1.4 ± 0.1	0.6 ± 0.1	< 0.4	< 0.3	< 0.33

**Table A-8e. Inductively Coupled Plasma-Mass Spectrometry Data from the Primary Adsorption Isotherm Experiment: Hanford Soil (4 of 4)**

Weight % of Hanford Soil	Gadolinium µg/L	Dysprosium µg/L	Holmium µg/L	Erbium µg/L	Ytterbium µg/L
100 %	<0.02	<0.02	<0.02	<0.02	<0.02
100 %	<0.02	<0.02	<0.02	<0.02	<0.02
100 %	<0.02	<0.02	<0.02	<0.02	<0.02
100 %	<0.02	0.02 ± 0.01	<0.02	<0.02	<0.02
100 %	0.38 ± 0.05	<0.02	<0.02	<0.02	<0.02
100 %	1.03 ± 0.19	0.05 ± 0.05	0.03 ± 0.01	0.06 ± 0.01	0.06 ± 0.07
100 %	0.26 ± 0.02	0.03 ± 0.00	<0.02	<0.02	<0.02
100 %	0.22 ± 0.04	0.04 ± 0.01	<0.02	<0.02	<0.02
100 %	<0.02	2.01 ± 0.03	0.04 ± 0.01	<0.02	<0.02

Avg. ± Std. Dev. 0.47 ± 0.38 0.43 ± 0.88 0.03 ± 0.00

Hanford Soil	µmol/L	µmol/L	µmol/L	µmol/L	µmol/L
100 %	< 0.1	< 0.12	< 0.12	< 0.12	< 0.12
100 %	< 0.1	< 0.12	< 0.12	< 0.12	< 0.12
100 %	< 0.1	< 0.12	< 0.12	< 0.12	< 0.12
100 %	< 0.1	0.15 ± 0.04	< 0.12	< 0.12	< 0.12
100 %	2.4 ± 0.3	< 0.12	< 0.12	< 0.12	< 0.12
100 %	6.6 ± 1.2	0.33 ± 0.28	0.20 ± 0.04	0.4 ± 0.0	0.34 ± 0.41
100 %	1.7 ± 0.1	0.19 ± 0.02	< 0.12	< 0.12	< 0.12
100 %	1.4 ± 0.2	0.23 ± 0.04	< 0.12	< 0.12	< 0.12
100 %	< 0.1	12.39 ± 0.17	0.22 ± 0.03	< 0.12	< 0.12

**APPENDIX B.**

**Experiment B: Kinetics of Complexation**

BHI-00864  
Rev. 0

**Table B-1. Liquid Scintillation Counting of Strontium-90, pH, and Conductivity Data from the Kinetics Experiment**

	Time hours	Date of counting	Sr-90 Bq/mL	pCi/L	pH	Conductivity microSi/cm
<b>NC Apatite 1%</b>	1	12/27/95	0.42	11,349	7.88	147
	3	12/27/95	0.40	10,941	8.31	154
	5	12/27/95	0.44	11,882	7.73	140
	10	12/27/95	0.47	12,613	7.79	149
	15	12/27/95	0.52	13,982	6.81	150
	24	12/27/95	0.51	13,845	8.09	155
<b>Bone Char 5%</b>	1	12/27/95	0.29	7,958	9.23	197
	3	12/27/95	0.28	7,488	9.47	195
	5	12/27/95	0.30	8,163	9.26	195
	10	12/27/95	0.24	6,573	9.45	187
	15	12/28/95	0.21	5,619	9.63	200
	24	12/28/95	0.20	5,367	9.50	195
<b>Hydroxyapatite 5%</b>	1	12/28/95	0.33	8,804	7.96	171
	3	12/28/95	0.35	9,518	8.45	168
	5	12/28/95	0.34	9,245	8.38	157
	10	12/28/95	0.42	11,300	7.63	171
	15	12/28/95	0.42	11,305	7.62	171
	24	12/28/95	0.42	11,268	7.93	171
<b>Clinoptilolite 5%</b>	1	12/28/95	0.19	5,022	8.54	162
	3	12/28/95	0.17	4,669	8.49	163
	5	12/28/95	0.12	3,201	8.81	171
	10	12/28/95	0.06	1,711	8.79	176
	15	12/29/95	0.07	1,988	8.92	183
	24	12/29/95	0.06	1,535	9.34	193
<b>Soil</b>	1	12/29/95	0.38	10,202	8.27	147
	3	12/29/95	0.40	10,800	8.35	150
	5	12/30/95	0.43	11,547	9.03	159
	10	12/30/95	0.51	13,683	9.10	162
	15	12/30/95	0.50	13,600	8.67	157
	24	12/30/95	0.51	13,672	8.36	156
<b>Hanford Water</b>			0	-	7.99	150

**Table B-2. Atomic Adsorbtion Spectrophotometry Data from the Kinetics Experiment**

	Time hours	Na ppm		Mg ppm		Ca ppm		StD%
<b>NC Apatite 1%</b>	1	17	±	1.2	6.8	4.2	±	0.06
	3	15	±	0.1	0.6	4.3	±	0.01
	5	16	±	0.9	5.7	4.2	±	0.01
	10	16	±	0.8	4.9	4.2	±	0.01
	15	17	±	0.3	1.7	4.2	±	0.04
	24	17	±	1.3	7.7	4.2	±	0.03
<b>Bone Char 5%</b>	1	32	±	0.5	1.4	5.9	±	0.03
	3	30	±	0.04	0.1	6.0	±	0.004
	5	32	±	0.6	1.8	6.2	±	0.06
	10	33	±	0.7	2.0	6.8	±	0.03
	15	35	±	0.7	2.0	7.6	±	0.02
	24	35	±	0.6	1.6	7.1	±	0.02
<b>Hydroxyapatite 5%</b>	1	17	±	0.3	1.9	5.3	±	0.03
	3	17	±	1.2	6.8	5.5	±	0.005
	5	16	±	0.6	3.6	5.4	±	0.05
	10	18	±	0.1	0.3	5.6	±	0.01
	15	16	±	0.8	4.9	5.6	±	0.01
	24	16	±	1.3	7.8	5.7	±	0.04
<b>Clinoptilolite 5%</b>	1	35	±	0.8	2.2	2.7	±	0.08
	3	35	±	0.5	1.3	2.7	±	0.02
	5	44	±	0.3	0.6	1.9	±	0.06
	10	51	±	0.4	0.8	1.2	±	0.01
	15	53	±	0.01	0.03	1.6	±	0.02
	24	57	±	0.4	0.7	1.6	±	0.01
<b>Soil</b>	1	16	±	0.3	1.9	4.3	±	0.03
	3	16	±	0.5	2.9	4.3	±	0.03
	5	16	±	0.2	1.4	4.2	±	0.05
	10	22	±	0.6	2.6	4.2	±	0.008
	15	16	±	0.5	3.1	4.2	±	0.01
	24	17	±	1.0	6.0	4.1	±	0.03
<b>Hanford Water</b>		16	±	0.1	0.3	4.6	±	0.01
						0.2		0.2
						20.8	±	0.8

**Table B-3. Inductively Coupled Plasma-Mass Spectrometry Data from the Kinetics Experiment (Page 1 of 7)**

	Time hour s	Chromium			Manganese			Iron		
		Value	Error	StD%	Value	Error	StD%	Value	Error	StD%
<b>NC Apatite 1%</b>	1	0.35 ±	0.07	19.7	7.6 ±	0	0.2	242 ±	1	0.4
	3	<0.2			11.5 ±	0.2	1.7	333 ±	3.6	1.1
	5	<0.2			9.7 ±	0.1	0.6	290 ±	4.5	1.6
	10	0.38 ±	0.03	7.4	8.1 ±	0.1	1	245 ±	5.9	2.4
	15	<0.2			8.4 ±	0.1	1.1	245 ±	8.6	3.5
	24	0.22 ±	0.01	6.6	8.6 ±	0.5	5.3	238 ±	13.5	5.7
<b>Bone Char 5%</b>	1	1.47 ±	0.04	2.8	2.2 ±	0	0.9	214 ±	8.6	4
	3	0.83 ±	0.02	2.5	5.2 ±	0.2	3	318 ±	9	2.8
	5	0.89 ±	0.02	2.1	4.1 ±	0	1	277 ±	2.9	1
	10	0.81 ±	0.03	3.5	0.98 ±	0	3.2	139 ±	3.6	2.6
	15	1.09 ±	0.01	1.3	1.51 ±	0	2.4	148 ±	2.2	1.5
	24	1.05 ±	0.03	2.5	1.67 ±	0	1.2	135 ±	3.8	2.9
<b>Hydroxyapatite 5%</b>	1	3.98 ±	0.01	0.3	1 ±	0	1.8	361 ±	3	0.8
	3	3.85 ±	0.03	0.7	1.14 ±	0	1.4	391 ±	1	0.2
	5	3.76 ±	0.08	2.1	1 ±	0	1.1	358 ±	7.3	2
	10	3.68 ±	0.07	1.9	1.29 ±	0	0.6	366 ±	7.5	2.1
	15	3.45 ±	0.02	0.5	1.35 ±	0	1.1	376 ±	14	3.7
	24	3.9 ±	0.01	0.3	1.56 ±	0	1.2	419 ±	4.8	1.1
<b>Clinoptilolite 5%</b>	1	0.58 ±	0.02	4	24.4 ±	0.2	0.6	677 ±	9.8	1.5
	3	0.94 ±	0.07	7.3	28.5 ±	0.1	0.5	779 ±	7.6	1
	5	0.7 ±	0.01	2	25.1 ±	0.3	1	755 ±	9.1	1.2
	10	0.76 ±	0.02	3.3	22.3 ±	0.1	0.5	700 ±	5.1	0.7
	15		na			na			na	
	24		na			na			na	
<b>Soil</b>	1	0.24 ±	0.02	7.4	6.6 ±	0	0.7	172 ±	5.5	3.2
	3	<0.2			8.5 ±	0.1	0.7	207 ±	2.2	1.1
	5	0.31 ±	0.01	3.4	6.9 ±	0.2	3.1	180 ±	0.3	0.2
	10	0.39 ±	0.02	5.9	10.3 ±	0.1	0.6	258 ±	2.3	0.9
	15	0.26 ±	0.01	3	9.3 ±	0.2	2	201 ±	3.7	1.9
	24	0.23 ±	0.01	3.6	7.3 ±	0.1	1.1	178 ±	2.5	1.4
<b>Hanford Water</b>		<0.2			0.34 ±	0	5.7	170 ±	15	8.8

**Table B-3. Inductively Coupled Plasma-Mass Spectrometry Data from the Kinetics Experiment (Page 2 of 7)**

	Time Hours	Copper			Zinc			Arsenic		
		Value	Error	StD%	Value	Error	StD%	Value	Error	StD%
<b>NC Apatite 1%</b>	1	3.85 ±	0.12	3.2	1.88 ±	0	1.2	0.18 ±	0.03	14.4
	3	2.86 ±	0.04	1.5	3.3 ±	0.1	2.3	0.2 ±	0.02	10.4
	5	2.24 ±	0.02	1	2.54 ±	0.1	3.9	0.22 ±	0.04	19.9
	10	4.13 ±	0.06	1.4	3.38 ±	0.1	2.4	0.33 ±	0.04	13.3
	15	3.15 ±	0.02	0.5	3.75 ±	0.1	1.3	0.36 ±	0.04	11.7
	24	2.32 ±	0.04	1.6	1.54 ±	0.2	10.4	0.37 ±	0.05	13.2
<b>Bone Char 5%</b>	1	5.01 ±	0.07	1.3	3.44 ±	0	1.2	0.14 ±	0.04	29.7
	3	3.62 ±	0.1	2.8	8.76 ±	0.2	2.2	0.15 ±	0.01	5.6
	5	4.16 ±	0.1	2.3	3.75 ±	0.1	3.4	0.16 ±	0.02	11.1
	10	2.17 ±	0.02	0.9	1.31 ±	0.1	3.6	0.17 ±	0.03	19.6
	15	1.57 ±	0.06	4.1	6.72 ±	0.1	1.6	0.12 ±	0.02	15.3
	24	1.26 ±	0.02	1.4	1.78 ±	0	2.3	0.18 ±	0.01	4.3
<b>Hydroxyapatite 5%</b>	1	1.02 ±	0.02	1.5	2.02 ±	0.2	8.4	0.05 ±	0	7
	3	1.34 ±	0.08	6.2	2.86 ±	0	0.9	0.06 ±	0.01	12.7
	5	1.08 ±	0.08	7.3	2.08 ±	0	0.7	0.06 ±	0.01	22.8
	10	0.88 ±	0.03	2.9	1.36 ±	0.1	6.4	0.08 ±	0.01	10.7
	15	0.88 ±	0.04	4.4	1.87 ±	0.1	5.2	0.1 ±	0.01	8.5
	24	0.86 ±	0.03	3.2	2.03 ±	0.1	5.9	0.09 ±	0.01	6.9
<b>Clinoptilolite 5%</b>	1	4.7 ±	0.14	3	3.08 ±	0.2	5.4	0.62 ±	0.03	4.3
	3	4.81 ±	0.05	1	3.84 ±	0.3	7.2	0.6 ±	0.02	3
	5	6.8 ±	0.14	2.1	7.71 ±	0.2	3.1	0.75 ±	0.03	4.6
	10	5.32 ±	0.04	0.7	8.86 ±	0.1	1.5	1.01 ±	0.05	5.2
	15	na			9.18	1.38				
	24	na			11.5	1.41				
<b>Soil</b>	1	2.31 ±	0.06	2.6	2.59 ±	0.1	3.4	0.11 ±	0.02	17.9
	3	3.68 ±	0.12	3.2	3.86 ±	0.2	4.2	0.12 ±	0.01	8
	5	2.78 ±	0.14	5	2.17 ±	0.2	6.8	0.16 ±	0.01	5.8
	10	2.52 ±	0.02	0.8	2.64 ±	0	0.4	0.2 ±	0.05	24.8
	15	2.31 ±	0.09	4	1.97 ±	0.1	4.3	0.23 ±	0.03	12.6
	24	2.59 ±	0.1	4	2.84 ±	0.2	7.1	0.25 ±	0.02	8.9
<b>Hanford Water</b>		1.44 ±	0.07	5	4.61 ±	0.2	4.1	<0.05		

**Table B-3. Inductively Coupled Plasma-Mass Spectrometry Data from the Kinetics Experiment (Page 3 of 7)**

	Time Hours	Strontium			Yttrium			Silver		
		Value	Error	StD%	Value	Error	StD%	Value	Error	StD%
<b>NC Apatite 1%</b>	1	56.8 ±	0.7	1.3	0.27 ±	0.1	19.9	<0.02		
	3	56 ±	0.9	1.6	1.03 ±	0.1	8.2	<0.02		
	5	60.8 ±	0.4	0.7	0.57 ±	0	5.4	<0.02		
	10	75.5 ±	1.1	1.4	2.33 ±	0.1	5.7	<0.02		
	15	73 ±	0.4	0.6	0.12 ±	0	7.5	<0.02		
	24	76.9 ±	0.8	1.1	0.06 ±	0	35.4	<0.02		
<b>Bone Char 5%</b>	1	41.2 ±	0.7	1.7	0.2 ±	0	15.9	<0.02		
	3	41.9 ±	0.6	1.5	0.57 ±	0.1	13.2	<0.02		
	5	42.8 ±	0.3	0.7	0.4 ±	0.1	12.2	<0.02		
	10	41.8 ±	0.5	1.1	0.07 ±	0	15.3	<0.02		
	15	38.3 ±	0.2	0.6	0.17 ±	0	12.9	<0.02		
	24	38.5 ±	0.4	1	0.15 ±	0	12.4	0.024 ±	0.01	40.1
<b>Hydroxyapatite 5%</b>	1	47.2 ±	0.4	0.9	0.08 ±	0	14.5	0.022 ±	0	12.2
	3	54.6 ±	0	0.1	<0.05			<0.02		
	5	50 ±	0.5	1	<0.05			<0.02		
	10	61.6 ±	0.5	0.8	<0.05			<0.02		
	15	60.7 ±	0.8	1.3	<0.05			<0.02		
	24	66.3 ±	0.6	0.9	<0.05			<0.02		
<b>Clinoptilolite 5%</b>	1	25.3 ±	0.2	1	2.37 ±	0.1	4.5	0.021 ±	0.01	27.3
	3	24.5 ±	0.4	1.4	2.6 ±	0.1	3.9	<0.02		
	5	11.8 ±	0.3	2.3	2 ±	0.1	5.3	0.036 ±	0.01	16.3
	10	7.86 ±	0.12	1.5	1.82 ±	0.1	5.9	<0.02		
	15	10.3			na			0.028		
	24	8.71			na			0.029		
<b>Soil</b>	1	52.5 ±	0.2	0.5	0.09 ±	0	14.6	<0.02		
	3	60.3 ±	0.6	1.1	0.28 ±	0	12	<0.02		
	5	59.3 ±	1.2	2.1	0.09 ±	0	20.8	<0.02		
	10	66.4 ±	0.8	1.2	0.44 ±	0.1	13	0.114 ±	0.03	23.8
	15	71.1 ±	0.5	0.7	0.15 ±	0	20	0.02 ±	0.01	40.6
	24	68.7 ±	1.6	2.3	0.06 ±	0	24.3	0.032 ±	0.01	20.8
<b>Hanford Water</b>		5.76 ±	0.16	2.9	0.06 ±	0	13.5	0.023 ±	0.01	20.9

**Table B-3. Inductively Coupled Plasma-Mass Spectrometry Data from the Kinetics Experiment (Page 4 of 7)**

	Time Hours	Cesium			Barium			Lanthanum		
		Value	Error	StD%	Value	Error	StD%	Value	Error	StD%
<b>NC Apatite 1%</b>	1	0.27 ± 0	0.01	2.6	10.5 ± 0	0.1	1.1	0.27 ± 0	0.01	2.6
	3	0.3 ± 0	0.01	2.5	11.5 ± 0	0.1	0.5	0.95 ± 0	0.05	5.1
	5	0.25 ± 0	0	1.7	11.4 ± 0	0.2	2.1	0.5 ± 0	0.15	29.1
	10	0.26 ± 0	0.01	2.6	11.6 ± 0	0.2	1.4	1.33 ± 0	0.14	10.7
	15	0.26 ± 0	0	1.4	11.8 ± 0	0.2	1.3	<0.1		
	24	0.25 ± 0	0.02	7.5	12.8 ± 0	0.2	1.9	<0.1		
<b>Bone Char 5%</b>	1	0.19 ± 0	0	1.5	11.7 ± 0	0.1	1	0.19 ± 0	0.01	7.1
	3	0.2 ± 0	0.01	5.1	12.4 ± 0	0.1	1	0.38 ± 0	0.02	4.7
	5	0.21 ± 0	0	0.8	12.4 ± 0	0.1	0.5	0.35 ± 0	0.02	5.3
	10	0.25 ± 0	0.01	3.1	10.7 ± 0	0.1	0.9	<0.1		
	15	0.19 ± 0	0	1.4	11.5 ± 0	0.1	0.8	0.14 ± 0	0.02	15.7
	24	0.19 ± 0	0	2	10.9 ± 0	0.1	0.8	0.14 ± 0	0.05	34.5
<b>Hydroxyapatite 5%</b>	1	0.18 ± 0	0.01	3	9.9 ± 0	0.1	1.2	0.25 ± 0	0.02	6.6
	3	0.2 ± 0	0	1.1	11.3 ± 0	0.1	0.9	<0.1		
	5	0.22 ± 0	0	1.7	10.7 ± 0	0.1	0.5	<0.1		
	10	0.21 ± 0	0	0.7	12.1 ± 0	0.1	0.8	<0.1		
	15	0.2 ± 0	0	1	12.3 ± 0	0	0.3	<0.1		
	24	0.22 ± 0	0	0.8	13.7 ± 0	0.1	0.9	<0.1		
<b>Clinoptilolite 5%</b>	1	0.24 ± 0	0	0.9	10.3 ± 0	0.1	0.9	2.94 ± 0	0.19	6.3
	3	0.24 ± 0	0	0.6	11 ± 0	0.2	1.4	3.27 ± 0	0.06	1.8
	5	0.23 ± 0	0	1	8.15 ± 0	0.2	2.2	2.5 ± 0	0.1	4.1
	10	0.26 ± 0	0	0.8	6.86 ± 0	0.2	2.5	2.25 ± 0	0.09	4
	15	0.25			na			na		
	24	0.24			na			na		
<b>Soil</b>	1	0.23 ± 0	0	1.6	9.9 ± 0	0.1	0.9	<0.1		
	3	0.25 ± 0	0.01	2.3	10.8 ± 0	0.2	1.9	0.17 ± 0	0.01	4.4
	5	0.23 ± 0	0	0.6	10.2 ± 0	0.1	0.6	<0.1		
	10	0.24 ± 0	0	0.9	12.5 ± 0	0.2	1.9	0.34 ± 0	0.02	7.3
	15	0.23 ± 0	0.01	2.3	11.3 ± 0	0.2	1.5	<0.1		
	24	0.24 ± 0	0.01	1.9	11.2 ± 0	0	0.2	<0.1		
<b>Hanford Water</b>		0.25 ± 0	0	0.7	0.25 ± 0	0	8	<0.1		

**Table B-3. Inductively Coupled Plasma-Mass Spectrometry Data  
from the Kinetics Experiment (Page 5 of 7)**

	Time Hours	Cerium			Praseodymium			Samarium		
		Value	Error	StD%	Value	Error	StD%	Value	Error	StD%
<b>NC Apatite 1%</b>	1	0.27 ± 0.02	7.6	<0.05				<0.05		
	3	1.32 ± 0.03	2.4	0.16 ± 0	14.2	0.19 ± 0	0.07	35		
	5	0.77 ± 0.04	4.8	0.09 ± 0	10.3	0.1 ± 0	0.05	49		
	10	1.24 ± 0.04	3.1	0.19 ± 0	4.8	0.14 ± 0	0.04	25		
	15	0.18 ± 0.03	19.1	<0.05				<0.05		
	24	<0.1		<0.05				<0.05		
<b>Bone Char 5%</b>	1	0.33 ± 0.05	14.9	<0.05				0.07 ± 0	0.11	150
	3	0.9 ± 0.08	8.4	0.12 ± 0	6	0.1 ± 0	0.09	92		
	5	0.7 ± 0.04	5.6	0.09 ± 0	37.2	0.11 ± 0	0.03	24		
	10	<0.1		<0.05				<0.05		
	15	0.28 ± 0.01	3.5	<0.05				0.05 ± 0	0.06	111
	24	0.29 ± 0.05	15.6	<0.05				<0.05		
<b>Hydroxyapatite 5%</b>	1	0.17 ± 0.02	9.2	<0.05				<0.05		
	3	<0.1		<0.05				<0.05		
	5	<0.1		<0.05				<0.05		
	10	<0.1		<0.05				<0.05		
	15	<0.1		<0.05				<0.05		
	24	<0.1		<0.05				0.05 ± 0	0.03	50
<b>Clinoptilolite 5%</b>	1	6.23 ± 0.25	4	0.62 ± 0.1	12	0.47 ± 0.1	0.15	31		
	3	7.59 ± 0.18	2.3	0.79 ± 0	4.4	0.67 ± 0	0.1	16		
	5	7.05 ± 0.16	2.3	0.56 ± 0	1.4	0.54 ± 0	0.08	16		
	10	6.53 ± 0.11	1.7	0.58 ± 0	5.1	0.38 ± 0	0.08	22		
	15	na			na			na		
	24	na			na			na		
<b>Soil</b>	1	<0.1		<0.05				<0.05		
	3	0.46 ± 0.01	1.9	0.07 ± 0	14.5	0.1 ± 0	0.03	31		
	5	0.17 ± 0.04	21.2	<0.05				0.09 ± 0	0.07	78
	10	1.06 ± 0.12	11.5	0.13 ± 0	22.3	0.07 ± 0	0.01	19		
	15	0.2 ± 0.03	15.8	<0.05				<0.05		
	24	<0.1		<0.05				0.06 ± 0	0.07	116
<b>Hanford Water</b>		<0.1		<0.05				0.08 ± 0	0.05	62

**Table B-3. Inductively Coupled Plasma-Mass Spectrometry Data from the Kinetics Experiment (Page 6 of 7)**

	Time Hours	Europium			Gadolinium			Dysprosium		
		Value	Error	StD%	Value	Error	StD%	Value	Error	StD%
<b>NC Apatite 1%</b>	1	<0.05			0.03 ± 0	77	<0.05			
	3	0.09 ± 0.01	11		0.21 ± 0.1	49	0.15 ± 0.04	30		
	5	<0.05			0.12 ± 0	34	0.11 ± 0.05	49		
	10	0.08 ± 0.01	11		0.28 ± 0.1	25	0.32 ± 0.05	16		
	15	<0.05			0.03 ± 0	120	<0.05			
	24	<0.05			0.05 ± 0	3	<0.05			
<b>Bone Char 5%</b>	1	<0.05			0.07 ± 0.1	78	0.1 ± 0.05	52		
	3	<0.05			0.19 ± 0.1	49	0.08 ± 0.08	99		
	5	<0.05			0.15 ± 0.1	60	0.1 ± 0.03	35		
	10	<0.05			0.08 ± 0	49	<0.05			
	15	<0.05			0.06 ± 0	70	0.06 ± 0.02	37		
	24	<0.05			0.08 ± 0.1	77	<0.05			
<b>Hydroxyapatite 5%</b>	1	<0.05			0.04 ± 0	36	<0.05			
	3	<0.05			<0.02		<0.05			
	5	<0.05			<0.02		<0.05			
	10	<0.05			0.02 ± 0	85	<0.05			
	15	<0.05			0.03 ± 0	149	<0.05			
	24	<0.05			0.05 ± 0	9	<0.05			
<b>Clinoptilolite 5%</b>	1	0.11 ± 0.03	29		0.38 ± 0.1	20	0.52 ± 0.12	23		
	3	0.14 ± 0.01	10		0.55 ± 0.1	21	0.57 ± 0.08	14		
	5	0.12 ± 0.02	16		0.51 ± 0.1	12	0.53 ± 0.01	2		
	10	0.11 ± 0.05	48		0.46 ± 0	6	0.49 ± 0.14	29		
	15	na			na		na			
	24	na			na		na			
<b>Soil</b>	1	<0.05			0.07 ± 0	59	0.09 ± 0.04	44		
	3	0.06 ± 0.02	41		0.08 ± 0	29	0.09 ± 0.06	65		
	5	0.09 ± 0.06	66		0.05 ± 0.1	99	0.06 ± 0.03	48		
	10	0.06 ± 0.04	76		0.12 ± 0	7	0.12 ± 0.08	70		
	15	<0.05			0.08 ± 0	18	0.07 ± 0.03	43		
	24	<0.05			0.05 ± 0	18	0.05 ± 0.04	83		
<b>Hanford Water</b>		<0.05			0.42 ± 0	9	0.03 ± 0.02	54		

**Table B-3. Inductively Coupled Plasma-Mass Spectrometry Data from the Kinetics Experiment (Page 7 of 7)**

	Time Hours	Holmium			Erbium			Lead		
		Value	Error	StD%	Value	Error	StD%	Value	Error	StD%
<b>NC Apatite 1%</b>	1	<0.02			<0.05			0.19 ±	0.02	9.7
	3	0.05 ±	0.01	18	0.1 ±	0	13	0.31 ±	0.01	2.9
	5	0.03 ±	0	13	<0.05			0.19 ±	0.01	5.7
	10	0.04 ±	0.01	31	0.1 ±	0	31	0.1 ±	0	0.6
	15	<0.02			<0.05			0.17 ±	0.03	14.5
	24	<0.02			<0.05			<0.1		
<b>Bone Char 5%</b>	1	<0.02			<0.05			0.16 ±	0.01	9
	3	<0.02			0.15 ±	0.1	34	0.3 ±	0.03	8.8
	5	0.03 ±	0.02	84	0.11 ±	0	36	0.2 ±	0.02	7.9
	10	<0.02			0.07 ±	0	30	<0.1		
	15	0.02 ±	0.01	24	0.06 ±	0	63	0.12 ±	0.01	11.4
	24	<0.02			0.07 ±	0	47	<0.1		
<b>Hydroxyapatite 5%</b>	1	<0.02			<0.05			0.17 ±	0.02	9.6
	3	<0.02			<0.05			0.17 ±	0.01	3.9
	5	<0.02			<0.05			0.13 ±	0.01	10.4
	10	<0.02			<0.05			<0.1		
	15	<0.02			<0.05			0.13 ±	0.01	8.9
	24	<0.02			<0.05			<0.1		
<b>Clinoptilolite 5%</b>	1	0.08 ±	0.02	19	0.36 ±	0.1	26	0.58 ±	0.02	3.3
	3	0.11 ±	0.02	14	0.47 ±	0.1	23	0.64 ±	0.03	5
	5	0.08 ±	0.02	24	0.33 ±	0.1	16	1.45 ±	0.02	1.1
	10	0.08 ±	0.02	23	0.36 ±	0	3	0.73 ±	0.01	1.3
	15				na					na
	24				na					na
<b>Soil</b>	1	<0.02			0.05 ±	0	86	0.14 ±	0.02	11.4
	3	0.02 ±	0.02	82	0.03 ±	0	71	0.38 ±	0.01	3.1
	5	0.03 ±	0.01	26	0.04 ±	0	88	0.19 ±	0.02	8.6
	10	0.04 ±	0.02	48	0.06 ±	0	53	0.25 ±	0.04	14.1
	15	<0.02			0.06 ±	0	36	0.14 ±	0.01	7.1
	24	<0.02			0.03 ±	0	41	0.11 ±	0	3.5
<b>Hanford Water</b>		<0.02			<0.02			0.31 ±	0.03	9.3

BHI-00864  
Rev. 0

**APPENDIX C.**

**Experiment C: Loading Capacity of Sorbents**

BHI-00864  
Rev. 0

**Table C-1. Results of the Loading Capacity Experiment**

System	Concentration of Sr in solution after 48 hours agitation.			Calculated concentration of Sr in solid phase after 48 hours agitation.			Std	Std% C(solid)/C(sol'n)	Kd
	C(sol'n) ppm	Std	Std% C(solid) ppm	Std	Std% C(solid)/C(sol'n)				
3 g NC Apatite and 0.01 g Sr(NO <sub>3</sub> ) <sub>2</sub> /30 mL water	51.7	± 0.72	1.4	80.9	±	3.80	4.7	4.7	1.57
3 g Bone Char and 0.01 g Sr(NO <sub>3</sub> ) <sub>2</sub> /30 mL water	0.36	± 0.05	13.5	132.3	±	18.8	14.2	14.2	371
3 g Hydroxyapatite and 0.01 g Sr(NO <sub>3</sub> ) <sub>2</sub> /30 mL water	16.3	± 0.49	3.0	116.3	±	6.29	5.4	7.11	
3 g Clinoptilolite and 0.01 g Sr(NO <sub>3</sub> ) <sub>2</sub> /30 mL water	0.23	± 0.00	0.7	132.4	±	6.02	4.5	4.5	585
Original solution 0.01 g Sr(NO <sub>3</sub> ) <sub>2</sub> /30 mL water	132.6	± 6.0	4.5	0.0	±	0.0	0.0	0.0	0.00
3 g NC Apatite and 0.1 g Sr(NO <sub>3</sub> ) <sub>2</sub> /30 mL water	996	± 42	4.2	265	±	19	7.3	7.3	0.27
3 g Bone Char and 0.1 g Sr(NO <sub>3</sub> ) <sub>2</sub> /30 mL water	261	± 16	6.2	1,000	±	86	8.6	8.6	3.83
3 g Hydroxyapatite and 0.1 g Sr(NO <sub>3</sub> ) <sub>2</sub> /30 mL water	589	± 17	2.9	672	±	44	6.6	6.6	1.14
3 g Clinoptilolite and 0.1 g Sr(NO <sub>3</sub> ) <sub>2</sub> /30 mL water	143	± 4.9	3.4	1,118	±	76.2	6.8	6.8	7.80
Original solution 0.1 g Sr(NO <sub>3</sub> ) <sub>2</sub> /30 mL water	1,261	± 75	5.9	0	±	0	0.0	0.0	0.00
3 g NC Apatite and 1 g Sr(NO <sub>3</sub> ) <sub>2</sub> /30 mL water	12,684	± 1,064	8.4	934	±	158	17.0	17.0	0.07
3 g Bone Char and 1 g Sr(NO <sub>3</sub> ) <sub>2</sub> /30 mL water	10,400	± 1,122	10.8	3,218	±	588	18.3	18.3	0.31
3 g Hydroxyapatite and 1 g Sr(NO <sub>3</sub> ) <sub>2</sub> /30 mL water	11,321	± 1,369	12.1	2,296	±	438	19.1	19.1	0.20
3 g Clinoptilolite and 1 g Sr(NO <sub>3</sub> ) <sub>2</sub> /30 mL water	9,632	± 1,255	13.0	3,985	±	784	19.7	19.7	0.41
Original solution 1 g Sr(NO <sub>3</sub> ) <sub>2</sub> /30 mL water	13,618	± 2,008	14.7	0	±	0	0.0	0.0	0.00

BHI-00864  
Rev. 0

**APPENDIX D.**

**Experiment D: pH Stability Test**

BHI-00864  
Rev. 0

**Table D-1. Ion Chromatography Data from the pH Stability Test, Saturation Step**

Treatment	Fluoride	Chloride	Nitrate	Phosphate	Sulfate
Number	ppm	ppm	ppm	ppm	ppm
<b>Hanford Water</b>	1 <0.1	<0.1	12.8	<0.5	19.1
	2 <0.1	<0.1	13.1	<0.5	18.7
	3 <0.1	<0.1	12.8	<0.5	18.6
<b>NC Apatite 1%</b>	1 0.98	25.3*	23.5	2.12	124.2
	2 0.95	2.26	35.7	0.87	143.0
	3 0.61	1.86	20.0	0.57	147.6
<b>Bone Char 5%</b>	1 not analyzed	52.1	52.0	0.61	93.5
	2 not analyzed	50.6	26.0	0.48	89.5
	3 not analyzed	39.5	12.6	0.61	85.0
<b>Hydroxyapatite 5%</b>	1 0.20	28.3	7.1	<0.5	6.62
	2 0.15	12.7	16.8	0.75	8.50
	3 0.15	8.67	12.1	0.92	7.53
<b>Clinoptilolite 5%</b>	1 1.05	40.4	23.3	2.19	70.8
	2 1.08	62.5	42.0	4.92	136.6
	3 2.17	66.5	57.3	3.58	137.3
<b>Maximal error of analysis</b>					
<b>in percent</b>					
	1.74	2.48	0.38	0.29	0.67

\* An error was made during the preparation of the samples in Treatment 1 when 0.01M KCl solution was mistakenly added instead of Hanford simulated groundwater. After discovery of this mistake, the samples were dried in an oven and then saturated with Hanford groundwater. This error could have increased concentration of chloride in the solutions.

**Table D-2. Strontium-90 Data by Liquid Scintillation for the pH Stability Test, Saturation Step**

	pH of added water	Treatment 1			Treatment 2			Treatment 3		
		Sr-90		Sr-90	Sr-90		Sr-90	Sr-90		
		Date of counting	Bq/mL pCi/L							
NC Apatite 1%	9	12/30/95	0.888	23,998	1/5/96	0.706	19,070	1/22/96	0.738	19,957
	8	12/30/95	0.843	22,794	1/5/96	0.845	22,851	1/22/96	0.859	23,227
	7	12/30/95	0.707	19,118	1/5/96	0.810	21,891	1/22/96	0.772	20,851
	6	12/30/95	0.805	21,743	1/5/96	0.972	26,260	1/22/96	0.786	21,241
	5	12/30/95	0.867	23,440	1/5/96	0.955	25,815	1/22/96	0.867	23,426
Bone Char 5%	9	12/30/95	0.093	2,519	1/5/96	0.065	1,766	1/22/96	0.056	1,518
	8	12/30/95	0.107	2,887	1/5/96	0.088	2,383	1/22/96	0.065	1,766
	7	12/30/95	0.065	1,765	1/6/96	0.066	1,792	1/22/96	0.050	1,363
	6	12/30/95	0.103	2,794	1/6/96	0.086	2,320	1/22/96	0.066	1,793
	5	12/31/95	0.092	2,484	1/6/96	0.078	2,121	1/23/96	0.041	1,104
Hydroxyapatite 5%	9	12/31/95	0.485	13,118	1/6/96	0.505	13,635	1/23/96	0.452	12,218
	8	12/31/95	0.573	15,498	1/6/96	0.584	15,796	1/23/96	0.443	11,982
	7	12/31/95	0.489	13,229	1/6/96	0.576	15,568	1/23/96	0.490	13,245
	6	12/31/95	0.554	14,978	1/6/96	0.634	17,145	1/23/96	0.518	14,000
	5	12/31/95	0.598	16,149	1/6/96	0.572	15,467	1/23/96	0.520	14,050
Clinoptilolite 5%	9	12/31/95	0.061	1,652	1/6/96	0.024	636	1/23/96	0.019	516
	8	12/31/95	0.043	1,154	1/6/96	0.054	1,471	1/23/96	0.019	502
	7	12/31/95	0.029	793	1/6/96	0.030	818	1/23/96	0.015	396
	6	12/31/95	0.031	839	1/6/96	0.043	1,175	1/23/96	0.016	439
	5	12/31/95	0.024	653	1/7/96	0.041	1,097	1/23/96	0.036	982
Soil	9	1/1/96	0.847	22,900	1/7/96	0.656	17,716	1/24/96	0.644	17,394
	8	1/1/96	0.783	21,158	1/7/96	0.628	16,968	1/24/96	0.718	19,414
	7	1/1/96	0.730	19,724	1/7/96	0.633	17,102	1/24/96	0.723	19,552
	6	1/1/96	0.573	15,489	1/7/96	0.718	19,409	1/24/96	0.751	20,286
	5	1/1/96	0.732	19,790	1/7/96	0.696	18,798	1/24/96	0.846	22,867
NC Apatite 1%	48 hr	1/1/96	0.529	14,304	1/8/96	0.406	10,976	1/24/96	0.406	10,968
Bone Char 5%	48 hr	1/1/96	0.222	5,997	1/8/96	0.109	2,959	1/24/96	0.235	6,345
Hydroxyapatite	48 hr	1/1/96	0.345	9,322	1/8/96	0.284	7,677	1/24/96	0.337	9,106
Clinoptilolite 5%	48 hr	1/2/96	0.048	1,307	1/8/96	0.031	847	1/25/96	0.039	1,054
Soil	48 hr	1/2/96	0.512	13,833	1/8/96	0.396	10,704	1/25/96	0.480	12,982
Hanford Water	9		0.000	-		0.000	-		0.000	-
	8		0.000	-		0.000	-		0.000	-
	7		0.000	-		0.000	-		0.000	-
	6		0.000	-		0.000	-		0.000	-
	5		0.000	-		0.000	-		0.000	-
Hanford Water	48 hr, Satur.		0.000	-		0.000	-		0.000	-
NC Apatite 1%	Satur.	1/2/96	1.535	41,498	1/8/96	1.639	44,306	1/25/96	1.387	37,486
Bone Char 5%	Satur.	1/2/96	0.198	5,341	1/8/96	0.147	3,984	1/25/96	0.145	3,917
Hydroxyapatite 5%	Satur.	1/2/96	0.794	21,463	1/8/96	0.656	17,717	1/25/96	0.330	8,910
Clinoptilolite 5%	Satur.	1/2/96	0.072	1,957	1/8/96	0.030	803	1/25/96	0.082	2,218

**Table D-3. Conductivity and pH Data by Liquid Scintillation for the pH Stability Test, Saturation Step**

	pH of added water	Treatment 1		Treatment 2		Treatment 3	
		pH	Cond. microSi/cm	pH	Cond. microSi/cm	pH	Cond. microSi/cm
<b>NC Apatite 1%</b>	9	8.62	203	7.87	192	8.02	169
	8	8.48	195	8.06	198	8.41	193
	7	7.81	187	7.92	198	7.98	210
	6	8.91	193	7.90	219	8.46	197
	5	7.86	192	7.05	204	8.23	206
<b>Bone Char 5%</b>	9	9.18	504	9.02	508	9.06	509
	8	9.11	459	9.12	523	9.25	501
	7	9.06	409	9.18	486	9.05	506
	6	9.09	466	8.95	488	9.37	585
	5	9.09	475	8.66	543	9.31	513
<b>Hydroxyapatite 5%</b>	9	8.07	192	8.47	190	8.51	187
	8	8.05	192	8.27	202	8.34	190
	7	7.74	189	8.32	227	8.74	217
	6	7.52	193	8.25	211	8.50	208
	5	7.75	215	8.37	235	8.48	222
<b>Clinoptilolite 5%</b>	9	8.97	423	9.06	356	9.60	396
	8	9.47	320	9.62	364	9.91	404
	7	9.81	358	9.65	400	9.49	406
	6	9.67	406	9.53	394	9.95	418
	5	9.91	410	8.52	371	10.17	417
<b>Soil</b>	9	8.13	196	9.28	222	8.35	146
	8	8.99	200	8.78	197	8.27	169
	7	9.53	194	8.43	198	8.39	197
	6	9.76	206	8.91	241	8.44	197
	5	9.41	208	9.14	233	8.37	198
<b>NC Apatite 1%</b>	48 hr	8.10	169	8.39	132	8.22	190
<b>Bone Char 5%</b>	48 hr	9.11	191	9.08	191	8.45	214
<b>HAP 5%</b>	48 hr	8.30	184	8.63	168	8.44	149
<b>Clinoptilolite 5%</b>	48 hr	9.39	208	9.56	194	9.28	192
<b>Soil</b>	48 hr	8.78	165	8.81	141	8.50	164
<b>Hanford Water</b>	9	8.90	163	9.16	116	8.99	106
	8	7.09	119	7.80	115	8.03	149
	7	8.00	145	6.99	108	7.02	159
	6	6.36	118	6.10	141	6.06	147
	5	5.04	118	5.14	128	5.01	154
<b>Hanford Water</b>	48 hr, Satur.	8.15	152	8.21	118	8.20	153
<b>NC Apatite 1%</b>	Satur.	8.20	388	7.71	348	7.00	306
<b>Bone Char 5%</b>	Satur.	8.44	1076	8.74	936	10.13	940
<b>HAP 5%</b>	Satur.	8.35	308	8.00	262	8.82	223
<b>Clinoptilolite 5%</b>	Satur.	8.61	364	8.76	668	7.24	691

**Table D-4a. Atomic Absorption Spectrophometry Data for the pH Stability Test: Sodium**

	pH of added water	Treatment 1		Treatment 2		Treatment 3	
		ppm	StD%	ppm	StD%	ppm	StD%
<b>NC Apatite 1%</b>	9	19 ± 0.2	0.8	20 ± 0.8	4.1	18 ± 0.1	0.5
	8	21 ± 0.1	0.6	20 ± 0.6	3.0	18 ± 0.8	4.3
	7	22 ± 1.6	7.1	20 ± 0.9	4.5	18 ± 0.1	0.3
	6	20 ± 0.5	2.4	19 ± 0.2	0.9	17 ± 0.7	4.1
	5	22 ± 1.2	5.2	20 ± 0.1	0.7	19 ± 0.3	1.8
<b>Bone Char 5%</b>	9	137 ± 0.4	0.3	138 ± 1.7	1.2	157 ± 2.5	1.6
	8	118 ± 0.2	0.2	147 ± 1.3	0.9	119 ± 1.9	1.6
	7	138 ± 0.9	0.7	135 ± 0.3	0.2	268 ± 47	17.6
	6	122 ± 1.3	1.1	141 ± 0.3	0.2	255 ± 40	15.5
	5	127 ± 1.3	1.1	144 ± 0.2	0.2	192 ± 22	11.5
<b>Hydroxyapatite 5%</b>	9	18 ± 0.4	2.0	20 ± 1.4	7.0	19 ± 0.5	2.4
	8	19 ± 0.7	3.8	21 ± 0.4	2.0	18 ± 0.1	0.6
	7	20 ± 0.02	0.1	20 ± 0.46	2.3	19 ± 0.85	4.5
	6	21 ± 0.4	1.9	19 ± 1.0	5.1	18 ± 0.3	1.8
	5	20 ± 0.8	3.9	19 ± 0.2	1.1	18 ± 0.0	0.3
<b>Clinoptilolite 5%</b>	9	131 ± 0.9	0.7	112 ± 0.7	0.6	113 ± 0.9	0.8
	8	101 ± 1.2	1.2	101 ± 0.4	0.4	109 ± 0.2	0.1
	7	110 ± 0.2	0.2	na	na	114 ± 0.2	0.1
	6	113 ± 0.8	0.7	105 ± 2.8	2.7	112 ± 1.3	1.2
	5	111 ± 1.0	0.9	110 ± 1.8	1.7	108 ± 0.2	0.2
<b>Soil</b>	9	21 ± 1.1	5.2	21 ± 2.3	10.9	18 ± 0.7	4.0
	8	22 ± 1.4	6.5	21 ± 0.3	1.2	18 ± 0.005	0.03
	7	21 ± 0.05	0.2	19 ± 1.10	5.8	17 ± 0.007	0.04
	6	19 ± 0.4	2.3	21 ± 0.9	4.1	16 ± 0.7	4.5
	5	20 ± 0.3	1.4	20 ± 1.7	8.4	17 ± 0.4	2.2
<b>NC Apatite 1%</b>	48 hr	16 ± 0.6	3.9	17 ± 2.6	14.9	13 ± 0.05	0.4
<b>Bone Char 5%</b>	48 hr	31 ± 0.2	0.5	34 ± 3.0	8.9	35 ± 0.5	1.5
<b>Hydroxyapatite 5%</b>	48 hr	17 ± 0.1	0.8	16 ± 1.7	10.5	14 ± 0.3	2.3
<b>Clinoptilolite 5%</b>	48 hr	55 ± 0.0	0.1	54 ± 2.2	4.0	55 ± 0.3	0.6
<b>Soil</b>	48 hr	17 ± 0.9	5.4	17 ± 2.5	14.8	14 ± 0.8	5.4
<b>Hanford Water</b>	9	15 ± 0.5	3.6	16 ± 2.6	16.1	14 ± 0.04	0.3
	8	16 ± 0.9	5.9	17 ± 2.7	16.4	16 ± 0.3	1.9
	7	16 ± 0.1	0.7	15 ± 0.8	5.4	14 ± 0.4	2.7
	6	17 ± 0.8	4.8	16 ± 0.2	1.6	15 ± 0.4	2.6
	5	16 ± 0.3	1.8	15 ± 0.1	0.8	16 ± 0.3	2.1
<b>Hanford Water</b>	48 hr, Satur.	15 ± 1.5	10.3	13 ± 0.1	1.0	14 ± 0.4	2.6
<b>NC Apatite 1%</b>	Satur.	30 ± 0.5	1.6	25 ± 0.1	0.4	24 ± 0.8	3.5
<b>Bone Char 5%</b>	Satur.	320 ± 51	15.9	336 ± 36	10.7	241 ± 8	3.4
<b>Hydroxyapatite 5%</b>	Satur.	25 ± 0.4	1.4	33 ± 0.4	1.2	22 ± 0.8	3.6
<b>Clinoptilolite 5%</b>	Satur.	107 ± 2	1.9	283 ± 66	23.3	189 ± 2	1.2

**Table D-4b. Atomic Absorption Spectrophotometry Data for the  
pH Stability Test: Magnesium**

	pH of added water	Treatment 1				Treatment 2				Treatment 3			
		ppm	StD%		ppm	StD%		ppm	StD%		ppm	StD%	
<b>NC Apatite 1%</b>	9	2.48 ± 0.004	0.2		2.24 ± 0.014	0.6		2.89 ± 0.02	0.8				
	8	2.05 ± 0.004	0.2		2.77 ± 0.020	0.7		3.46 ± 0.02	0.7				
	7	2.34 ± 0.029	1.2		2.56 ± 0.011	0.4		3.12 ± 0.02	0.7				
	6	1.97 ± 0.001	0.05		3.07 ± 0.022	0.7		3.02 ± 0.005	0.2				
	5	2.36 ± 0.002	0.1		2.95 ± 0.020	0.7		3.69 ± 0.02	0.7				
<b>Bone Char 5%</b>	9	1.27 ± 0.001	0.1		1.03 ± 0.023	2.3		1.21 ± 0.01	0.8				
	8	1.04 ± 0.002	0.2		1.52 ± 0.029	1.9		1.24 ± 0.01	0.7				
	7	1.27 ± 0.0005	0.04		1.38 ± 0.000	0.0		1.43 ± 0.01	0.6				
	6	1.38 ± 0.002	0.2		1.52 ± 0.028	1.8		1.43 ± 0.01	0.6				
	5	1.61 ± 0.005	0.3		1.38 ± 0.001	0.1		1.24 ± 0.01	0.6				
<b>Hydroxyapatite 5%</b>	9	5.33 ± 0.008	0.1		5.27 ± 0.064	1.2		4.92 ± 0.05	1.0				
	8	4.59 ± 0.005	0.1		6.02 ± 0.016	0.3		5.13 ± 0.02	0.4				
	7	5.01 ± 0.012	0.2		5.90 ± 0.013	0.2		6.04 ± 0.08	1.3				
	6	5.55 ± 0.021	0.4		6.03 ± 0.039	0.6		5.90 ± 0.06	1.0				
	5	5.89 ± 0.028	0.5		5.99 ± 0.037	0.6		6.26 ± 0.03	0.5				
<b>Clinoptilolite 5%</b>	9	0.27 ± 0.024	8.9		0.12 ± 0.014	11.0		0.20 ± 0.01	6.6				
	8	0.13 ± 0.004	3.3		0.16 ± 0.013	8.3		0.16 ± 0.01	8.3				
	7	0.13 ± 0.004	3.2			na		0.19 ± 0.01	7.0				
	6	0.13 ± 0.004	3.0		0.16 ± 0.022	13.9		0.12 ± 0.01	11.6				
	5	0.13 ± 0.004	2.9		0.15 ± 0.021	13.5		0.14 ± 0.01	9.2				
<b>Soil</b>	9	3.51 ± 0.015	0.4		2.13 ± 0.008	0.4		2.66 ± 0.01	0.5				
	8	2.84 ± 0.013	0.5		2.32 ± 0.010	0.4		3.09 ± 0.01	0.3				
	7	3.16 ± 0.015	0.5		2.11 ± 0.039	1.9		3.19 ± 0.02	0.6				
	6	2.91 ± 0.041	1.4		2.71 ± 0.042	1.5		3.32 ± 0.01	0.2				
	5	3.32 ± 0.015	0.5		2.58 ± 0.017	0.7		3.62 ± 0.003	0.1				
<b>NC Apatite 1%</b>	48 hr	4.51 ± 0.020	0.4		3.71 ± 0.022	0.6		3.83 ± 0.03	0.7				
<b>Bone Char 5%</b>	48 hr	6.76 ± 0.002	0.0		5.47 ± 0.057	1.1		6.52 ± 0.02	0.3				
<b>Hydroxyapatite 5%</b>	48 hr	5.60 ± 0.002	0.0		4.39 ± 0.031	0.7		4.43 ± 0.03	0.8				
<b>Clinoptilolite 5%</b>	48 hr	1.06 ± 0.022	2.0		0.76 ± 0.019	2.5		0.73 ± 0.01	1.6				
<b>Soil</b>	48 hr	4.45 ± 0.049	1.1		3.44 ± 0.036	1.1		3.80 ± 0.00	0.0				
<b>Hanford Water</b>	9	4.08 ± 0.020	0.5		4.52 ± 0.073	1.6		4.54 ± 0.03	0.6				
	8	4.35 ± 0.022	0.5		4.33 ± 0.075	1.7		4.44 ± 0.05	1.1				
	7	4.35 ± 0.023	0.5		4.46 ± 0.007	0.2		4.48 ± 0.01	0.2				
	6	4.37 ± 0.004	0.1		4.44 ± 0.020	0.5		4.44 ± 0.04	0.9				
	5	4.33 ± 0.051	1.2		4.41 ± 0.032	0.7		4.48 ± 0.03	0.8				
<b>Hanford Water</b>	48 hr, Satur.	4.62 ± 0.026	0.6		4.61 ± 0.006	0.1		4.59 ± 0.03	0.6				
<b>NC Apatite 1%</b>	Satur.	5.39 ± 0.062	1.1		6.35 ± 0.015	0.2		6.13 ± 0.02	0.3				
<b>Bone Char 5%</b>	Satur.	5.23 ± 0.004	0.1		4.73 ± 0.005	0.1		3.72 ± 0.03	0.8				
<b>Hydroxyapatite 5%</b>	Satur.	9.05 ± 0.109	1.2		7.92 ± 0.006	0.1		5.62 ± 0.02	0.3				
<b>Clinoptilolite 5%</b>	Satur.	1.25 ± 0.001	0.1		0.74 ± 0.012	1.7		0.79 ± 0.01	1.0				

**Table D-4c. Atomic Absorption Spectrophotometry Data for the pH Stability Test: Calcium**

	pH of added water	Treatment 1		Treatment 2		Treatment 3		StD%	StD%
		ppm	StD%	ppm	StD%	ppm	StD%		
NC Apatite 1%	9	32.2	± 0.5	1.6	26.7	± 0.5	1.8	28.9	± 0.8
	8	26.0	± 0.4	1.7	30.1	± 0.7	2.2	31.2	± 0.6
	7	28.5	± 0.5	1.7	29.9	± 0.4	1.5	29.5	± 0.2
	6	27.2	± 0.5	1.7	33.9	± 0.4	1.3	28.9	± 0.04
	5	30.4	± 0.5	1.6	11.1	± 0.4	3.2	50.8	± 0.4
Bone Char 5%	9	1.91	± 0.13	7.0	1.78	± 0.01	0.6	1.98	± 0.04
	8	1.80	± 0.02	1.2	2.28	± 0.01	0.6	1.86	± 0.01
	7	2.03	± 0.01	0.5	2.20	± 0.01	0.6	2.51	± 0.02
	6	2.26	± 0.002	0.1	1.87	± 0.01	0.8	2.19	± 0.02
	5	2.26	± 0.01	0.5	5.61	± 0.05	0.8	2.35	± 0.02
Hydroxyapatite 5%	9	31.1	± 0.01	0.04	27.8	± 0.2	0.8	24.6	± 0.1
	8	26.8	± 0.3	1.1	30.5	± 0.1	0.2	24.0	± 0.7
	7	28.6	± 0.01	0.1	29.8	± 0.4	1.3	28.7	± 0.4
	6	30.4	± 0.2	0.5	30.9	± 0.2	0.5	28.7	± 0.03
	5	33.1	± 0.03	0.1	10.2	± 0.1	0.8	30.4	± 0.4
Clinoptilolite 5%	9	2.11	± 0.02	1.0	1.42	± 0.03	2.1	1.53	± 0.08
	8	1.22	± 0.03	2.4	1.38	± 0.03	2.3	1.36	± 0.08
	7	1.22	± 0.03	2.6			na	1.66	± 0.02
	6	1.38	± 0.04	2.6	1.37	± 0.01	0.9	1.15	± 0.02
	5	1.18	± 0.09	7.8	2.12	± 0.02	0.8	1.15	± 0.02
Soil	9	30.2	± 0.4	1.5	27.9	± 0.2	0.5	21.5	± 0.7
	8	23.4	± 0.1	0.5	31.1	± 0.1	0.2	25.0	± 0.0
	7	25.7	± 0.7	2.6	28.8	± 0.1	0.2	26.9	± 0.2
	6	22.4	± 0.6	2.7	34.2	± 0.3	0.8	27.0	± 0.4
	5	26.3	± 0.2	0.6	32.7	± 0.6	1.9	29.7	± 0.3
NC Apatite 1%	48 hr	25.8	± 0.0	0.1	16.6	± 0.2	1.2	27.8	± 0.1
Bone Char 5%	48 hr	9.1	± 0.2	2.5	6.1	± 0.1	2.3	14.0	± 0.1
Hydroxyapatite 5%	48 hr	25.6	± 0.4	1.5	12.0	± 8.6	71.6	21.3	± 0.3
Clinoptilolite 5%	48 hr	2.82	± 0.03	1.1	1.81	± 0.01	0.5	2.26	± 0.01
Soil	48 hr	25.9	± 0.4	1.5	17.3	± 0.1	0.4	23.5	± 0.8
Hanford Water	9	26.4	± 0.1	0.2	11.6	± 0.2	1.8	21.1	± 4.5
	8	10.9	± 0.1	0.7	14.0	± 0.1	0.4	20.4	± 0.5
	7	19.3	± 0.1	0.3	10.9	± 0.1	1.2	14.7	± 0.4
	6	11.4	± 0.1	0.8	16.1	± 0.05	0.3	14.4	± 0.2
	5	11.7	± 0.1	0.8	12.4	± 0.1	1.2	17.1	± 0.4
Hanford Water	48hr, Satur.	88.2	± 3.9	4.5	27.1	± 0.2	0.7	30.3	± 0.7
NC Apatite 1%	Satur.	64.6	± 0.9	1.4	63.6	± 0.7	1.1	53.9	± 0.8
Bone Char 5%	Satur.	8.5	± 0.1	0.7	6.0	± 0.04	0.7	3.99	± 0.23
Hydroxyapatite 5%	Satur.	14.8	± 0.2	1.1	36.2	± 0.8	2.1	24.9	± 0.2
Clinoptilolite 5%	Satur.	17.4	± 0.2	1.1	6.0	± 0.1	1.9	4.80	± 0.09

**Table D-5a. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: NC Apatite (Page 1 of 4)**

of added water	pH	Chromium		Manganese			Iron		
	ppb	StD%	ppb	StD%	ppb	StD%	ppb	StD%	ppb
<b>Treatment 1</b>	9	7.4 ± 0.05	0.7	2.97 ± 0.1	3.3	365 ± 6.2	1.7		
	8	6.3		1.42		300			
	7	7.2		3.68		277			
	6	5.2		1.17		303			
	5	7.4		1.45		364			
	48 hr	0.4 ± 0.02	5.2	11.4 ± 0.13	1.2	302 ± 3	1		
	Sat.	50.1 ± 0.61	1.2	4.76 ± 0	0.1	847 ± 10.2	1.2		
	9	17.3		1.65		322			
	8	15.3 ± 0.13	0.9	1.61 ± 0.05	3	355 ± 9.4	2.7		
	7	14.1		1.28		336			
<b>Treatment 2</b>	6	15.1 ± 0.22	1.5	1.84 ± 0.02	0.9	405 ± 18.5	4.6		
	5	15.9		1.83		413			
	48 hr	6.3 ± 0.03	0.5	6.92 ± 0.05	0.7	277 ± 0.7	0.3		
	Sat.	54 ± 0.77	1.4	7.98 ± 0.11	1.4	925 ± 60.4	6.5		
	9	2.3		1.8		312			
	8	1.5 ± 0.02	1.1	1.62 ± 0.01	0.5	341 ± 3.7	1.1		
	7	2.4		1.95		326			
	6	4.3		1.42		338			
	5	2.4 ± 0.02	0.7	1.8 ± 0.03	1.4	388 ± 7.4	1.9		
	48 hr	1.3 ± 0.03	2.2	3.85 ± 0.04	1.1	322 ± 8.9	2.8		
<b>Treatment 3</b>	Sat.	6.6 ± 0.06	0.9	3.32 ± 0.04	1.2	617 ± 22	3.6		
	<b>Copper</b>			<b>Zinc</b>			<b>Arsenic</b>		
	9	4.75 ± 0.02	0.3	3.7 ± 0.06	1.7	1.92 ± 0.06	3.1		
	8	3.15		17.8		1.63			
	7	4.72		6.34		1.77			
	6	3.06		16		1.42			
	5	3.06		2.06		1.89			
	48 hr	1.69 ± 0.06	3.8	11.2 ± 0.18	1.6	0.31 ± 0.03	10.4		
	Sat.	17.7 ± 0.23	1.3	26.7 ± 0.19	0.7	2.85 ± 0.02	0.8		
	9	5.67		28.1		2.75			
<b>Treatment 2</b>	8	3.38 ± 0.03	1	5.33 ± 0.12	2.3	1.93 ± 0.09	4.6		
	7	2.38		3.33		1.96			
	6	2.56 ± 0.1	3.9	8.72 ± 0.23	2.6	1.82 ± 0.08	4.3		
	5	3		12.3		2.14			
	48 hr	5.12 ± 0.07	1.4	35.4 ± 0.62	1.8	0.53 ± 0.04	6.9		
	Sat.	8.15 ± 0.28	3.4	18.6 ± 0.15	0.8	2.24 ± 0.08	3.7		
	9	5.46		10		1.73			
	8	3.5 ± 0.02	0.6	4.99 ± 0.1	2	1.58 ± 0.06	3.5		
	7	3.3		11		1.65			
	6	3.35		10.9		1.34			
<b>Treatment 3</b>	5	2.78 ± 0.15	5.3	14.2 ± 0.19	1.3	1.39 ± 0.01	0.4		
	48 hr	2.09 ± 0.04	1.8	10.4 ± 0.15	1.4	0.49 ± 0.06	12.7		
	Sat.	4.11 ± 0.08	1.9	12.5 ± 0.2	1.6	1.95 ± 0.05	2.3		

**Table D-5a. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: NC Apatite (Page 2 of 4)**

of added water	pH	Strontium			Yttrium			Silver		
	ppb	ppb	StD%	ppb	ppb	StD%	ppb	ppb	StD%	
<b>Treatment 1</b>	9	142	± 2.9	2	0.28	± 0.01	5.3	0.072	± 0.01	16.5
	8	119			na		<0.02			
	7	109			0.33		<0.02			
	6	117			na		<0.02			
	5	129			na		<0.02			
	48 hr	76.2	± 0.5	0.6	0.19	± 0.01	5.5	<0.02		
	Sat.	255	± 2.6	1	na		<0.02			
<b>Treatment 2</b>	9	109			na		<0.02			
	8	127	± 1.5	1.2	na		<0.02			
	7	124			na		<0.02			
	6	142	± 2.5	1.8	na		<0.02			
	5	136			na		<0.02			
	48 hr	62.3	± 0.5	0.8	1.17	± 0.08	6.9	<0.02		
	Sat.	264	± 4.3	1.6	na		<0.02			
<b>Treatment 3</b>	9	116			na		<0.02			
	8	128	± 0.7	0.6	na		<0.02			
	7	124			na		0.025			
	6	119			na		<0.02			
	5	135	± 1.5	1.1	na		<0.02			
	48 hr	85	± 1.4	1.7	0.44	± 0.03	7.3	<0.02		
	Sat.	221	± 1.9	0.9	na		<0.02			
<b>Cesium</b>										
<b>Treatment 1</b>	9	0.221	± 0	0.6	25.1	± 0.22	0.9	0.47	± 0.02	3.9
	8	0.21			23.4		<0.1			
	7	0.216			19.9		0.12			
	6	0.222			22.2		<0.1			
	5	0.201			24.9		<0.1			
	48 hr	0.204	± 0.01	3.4	13.3	± 0.31	2.4	<0.1		
	Sat.	0.276	± 0	0.9	30.7	± 0.22	0.7	0.62	± 0.09	14
<b>Treatment 2</b>	9	0.193			13.9		<0.1			
	8	0.191	± 0	1.3	19.9	± 0.34	1.7	<0.1		
	7	0.184	± 0	0	20		<0.1			
	6	0.194	± 0.01	2.8	23.5	± 0.29	1.2	<0.1		
	5	0.185			23.7		0.23			
	48 hr	0.253	± 0	0.8	10	± 0.18	1.8	0.27	± 0.02	7
	Sat.	0.195	± 0.01	3.6	43.7	± 0.74	1.7	0.68	± 0.08	11.4
<b>Treatment 3</b>	9	0.489			21.7		<0.1			
	8	0.222	± 0	0.4	23	± 0.38	1.7	<0.1		
	7	0.263			22.6		<0.1			
	6	0.205			23.6		<0.1			
	5	0.192	± 0	1.7	27.2	± 0.22	0.8	<0.1		
	48 hr	0.206	± 0.01	3.5	17	± 0.02	0.1	<0.1		
	Sat.	0.196	± 0	0.2	43.3	± 0.33	0.8	<0.1		

**Table D-5a. Inductively Coupled Plasma-Mass Spectrometry Data for the  
pH Stability Test: NC Apatite (Page 3 of 4)**

	<b>pH</b> of added water	<b>Cerium</b>				<b>Praeseodymium</b>				<b>Samarium</b>			
		<b>ppb</b>	<b>StD%</b>	<b>ppb</b>	<b>StD%</b>	<b>ppb</b>	<b>StD%</b>	<b>ppb</b>	<b>StD%</b>	<b>ppb</b>	<b>StD%</b>	<b>ppb</b>	<b>StD%</b>
<b>Treatment 1</b>	9	0.25	± 0.01	2		0.07	± 0	4.5		0.1	± 0.03	33	
	8	<0.1				<0.05				<0.05			
	7	0.31				<0.05				0.09			
	6	<0.1				<0.05				<0.05			
	5	<0.1				<0.05				<0.05			
	48 hr	0.17	± 0.03	15.6		<0.05				<0.05			
	Sat.	0.87	± 0.05	6		0.1	± 0.01	12.9		0.08	± 0.11	133	
<b>Treatment 2</b>	9	0.1				<0.05				0.07			
	8	0.16	± 0.02	15.1		<0.05				<0.05			
	7	0.1				<0.05				<0.05			
	6	0.12	± 0.02	12.3		<0.05				0.08	± 0.07	88	
	5	0.28				<0.05				<0.05			
	48 hr	0.38	± 0.03	7.7		<0.05				<0.05			
	Sat.	1.35	± 0.05	3.6		0.24	± 0.01	5		0.11	± 0.04	36	
<b>Treatment 3</b>	9	0.13				<0.05				<0.05			
	8	<0.1				<0.05				0.06	± 0.05	87	
	7	<0.1				<0.05				<0.05			
	6	0.18				<0.05				0.09			
	5	<0.1				<0.05				<0.05			
	48 hr	0.15	± 0	1.9		<0.05				<0.05			
	Sat.	<0.1				<0.05				0.07	± 0.08	111	
<b>Treatment 1</b>	<b>Europium</b>				<b>Gadolinium</b>				<b>Dysprosium</b>				
	9	0.061	± 0.04	73		0.08	± 0.01	14		0.11	± 0.02	23	
	8	0.083				0.04				0.13			
	7	0.067				<0.02				<0.05			
	6	<0.05				<0.02				<0.05			
	5	0.092				0.03				0.09			
	48 hr	<0.05				0.05	± 0.01	25		<0.05			
	Sat.	0.097	± 0.02	23		0.09	± 0.04	43		0.32	± 0.12	38	
<b>Treatment 2</b>	9	<0.05				<0.02				0.06			
	8	0.054	± 0.03	62		0.06	± 0.03	42		<0.05			
	7	<0.05				0.03				<0.05			
	6	0.055	± 0.04	72		0.03	± 0.03	92		<0.05			
	5	<0.05				0.04				0.25			
	48 hr	0.055	± 0.01	10		0.06	± 0.04	58		1.28	± 0.19	15	
	Sat.	0.143	± 0.01	6		0.2	± 0.01	7		0.89	± 0.14	16	
<b>Treatment 3</b>	9	0.065				0.06				0.19			
	8	0.053	± 0.03	63		0.05	± 0.01	16		<0.05			
	7	<0.05				<0.02				0.08			
	6	<0.05				0.04				0.06			
	5	0.051	± 0.03	49		0.05	± 0.04	72		0.14	± 0.08	56	
	48 hr	<0.05				0.09	± 0.05	51		0.43	± 0.04	10	
	Sat.	0.07	± 0.01	7		0.03	± 0.05	151		0.13	± 0.11	83	

**Table D-5a. Inductively Coupled Plasma-Mass Spectrometry Data for the  
pH Stability Test: NC Apatite (Page 4 of 4)**

	pH of added water	ppb	Holmium StD%	ppb	Erbium StD%	ppb	Lead StD%
<b>Treatment 1</b>	9	<0.02		<0.05		<0.1	
	8	<0.02		<0.05		<0.1	
	7	<0.02		<0.05		<0.1	
	6	<0.02		0.06		0.11	
	5	<0.02		<0.05		<0.1	
	48 hr	<0.02		<0.05		0.14 ± 0.01	5.4
	Sat.	0.028 ± 0.02	75	<0.05		0.73 ± 0.02	2
<b>Treatment 2</b>	9	<0.02		<0.05		0.34	
	8	<0.02		<0.05		<0.1	
	7	<0.02		<0.05		0.13	
	6	<0.02		0.06 ± 0.03	45	<0.1	
	5	0.041		0.06		0.23	
	48 hr	0.07 ± 0.01	18	<0.05		2.03 ± 0.08	4.2
	Sat.	0.056 ± 0.01	20	0.06 ± 0.03	52	0.92 ± 0.01	1.2
<b>Treatment 3</b>	9	<0.02		<0.05		0.17	
	8	<0.02		<0.05		<0.1	
	7	<0.02		<0.05		0.19	
	6	<0.02		<0.05		0.32	
	5	<0.02		<0.05		0.18 ± 0.03	16.5
	48 hr	<0.02		<0.05		0.5 ± 0.01	1.2
	Sat.	<0.02		0.08 ± 0.06	72	0.15 ± 0.03	20.3

**Table D-5b. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Bone Char (Page 1 of 4)**

of added water	pH	Chromium		Manganese		Iron	
	ppb	StD%	ppb	StD%	ppb	StD%	
<b>Treatment 1</b>	9	11.4 ± 0.21	1.8	8.21 ± 0.05	0.6	197 ± 3.7	1.9
	8	10.5 ± 0.09	0.9	0.85 ± 0.02	2.1	17 ± 1.2	6.9
	7	8.0 ± 0.08	1.0	1.69 ± 0.01	0.7	59 ± 0.9	1.5
	6	7.7 ± 0.12	1.5	3.25 ± 0.04	1.2	78 ± 2.2	2.9
	5	7.2 ± 0.20	2.7	0.42 ± 0.00	0.8	12 ± 2.4	20.1
	48 hr	0.7 ± 0.01	1.2	0.77 ± 0.01	1.6	103 ± 0.7	0.6
	Satur	63.4 ± 1.0	1.6	2.72 ± 0.01	0.5	120 ± 5.4	4.5
<b>Treatment 2</b>	9	19.5 ± 0.19	1.0	0.64 ± 0.02	2.7	13 ± 0.8	6.3
	8	14.2 ± 0.22	1.5	0.93 ± 0.01	1.2	41 ± 0.8	1.9
	7	12.4 ± 0.18	1.5	0.87 ± 0.01	1.1	17 ± 2.2	12.4
	6	12.1 ± 0.15	1.3	0.51 ± 0.01	1.8	17 ± 1.8	10.2
	5	15.6 ± 0.13	0.8	0.74 ± 0.02	2.3	11 ± 1.7	15.2
	48 hr	1.6 ± 0.05	3.1	1.58 ± 0.02	1.3	69 ± 2.1	3.0
	Sat.	28.8 ± 0.33	1.1	2.20 ± 0.02	1.1	137 ± 2.5	1.8
<b>Treatment 3</b>	9	6.5 ± 0.10	1.5	0.68 ± 0.04	6.1	<10	
	8	7.1 ± 0.07	0.9	0.71 ± 0.00	0.6	19 ± 1.0	5.4
	7	6.2 ± 0.05	0.9	0.57 ± 0.02	3.3	14 ± 1.9	14.1
	6	7.8 ± 0.11	1.4	0.63 ± 0.01	2.3	17 ± 3.3	19.5
	5	7.3 ± 0.07	1.0	0.63 ± 0.03	4.8	26 ± 1.5	5.8
	48 hr	0.9 ± 0.03	3.6	1.55 ± 0.01	0.7	189 ± 1.0	0.5
	Sat.	15.8 ± 0.04	0.3	0.76 ± 0.01	1.4	55 ± 0.5	0.8
		Copper	Zinc		Arsenic		
<b>Treatment 1</b>	9	8.20 ± 0.07	0.8	1.97 ± 0.22	10.9	2.09 ± 0.04	2.1
	8	6.71 ± 0.16	2.4	19.9 ± 0.20	1.0	1.83 ± 0.05	2.9
	7	5.55 ± 0.14	2.5	6.19 ± 0.27	4.4	1.55 ± 0.06	3.6
	6	3.09 ± 0.06	1.8	4.18 ± 0.11	2.7	1.30 ± 0.06	4.6
	5	2.62 ± 0.05	2.0	1.34 ± 0.12	8.7	1.26 ± 0.05	4.3
	48 hr	1.34 ± 0.08	6.0	6.13 ± 0.14	2.3	0.17 ± 0.03	15.8
	Sat.	6.32 ± 0.11	1.7	2.47 ± 0.06	2.5	3.03 ± 0.08	2.6
<b>Treatment 2</b>	9	3.87 ± 0.08	2.1	20.3 ± 0.18	0.9	1.91 ± 0.02	0.9
	8	2.55 ± 0.33	12.8	7.29 ± 0.10	1.4	1.60 ± 0.05	2.9
	7	1.79 ± 0.06	3.4	3.81 ± 0.18	4.6	1.31 ± 0.01	0.6
	6	1.28 ± 0.02	1.2	2.47 ± 0.06	2.5	1.35 ± 0.10	7.2
	5	1.82 ± 0.06	3.4	6.36 ± 0.13	2.0	1.44 ± 0.07	4.6
	48 hr	0.99 ± 0.04	3.7	6.81 ± 0.11	1.6	0.30 ± 0.01	4.9
	Sat.	2.07 ± 0.02	0.9	12.2 ± 0.23	1.9	1.47 ± 0.04	2.7
<b>Treatment 3</b>	9	1.84 ± 0.12	6.5	2.74 ± 0.14	5.2	1.19 ± 0.04	3.6
	8	1.41 ± 0.06	4.2	7.11 ± 0.11	1.6	0.98 ± 0.05	4.7
	7	0.75 ± 0.05	6.4	1.37 ± 0.05	3.7	0.96 ± 0.02	1.6
	6	0.79 ± 0.02	1.9	4.02 ± 0.03	0.8	1.03 ± 0.01	1.1
	5	1.43 ± 0.01	1.0	6.09 ± 0.16	2.6	0.86 ± 0.05	5.6
	48 hr	1.32 ± 0.05	3.9	5.43 ± 0.18	3.4	0.17 ± 0.01	5.7
	Sat.	0.77 ± 0.06	7.8	1.56 ± 0.25	15.8	1.03 ± 0.06	5.8

**Table D-5b. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Bone Char (Page 2 of 4)**

	pH of added water	Strontium			Yttrium			Silver		
		ppb	StD%	ppb	StD%	ppb	StD%	ppb	StD%	ppb
<b>Treatment 1</b>	9	12.3 ± 0.23	1.9	0.91 ± 0.04	4.9	0.452 ± 0.00	1.6			
	8	12.5 ± 0.12	0.9	1.41 ± 0.14	10.2	0.282 ± 0.03	12.2			
	7	10.8 ± 0.12	1.1	338 ± 4	1.3	0.205 ± 0.00	3.7			
	6	14.6 ± 0.07	0.5	1.15 ± 0.15	13.3	0.160 ± 0.01	6.2			
	5	14.8 ± 0.28	1.9	81.7 ± 1.2	1.5	0.137 ± 0.02	18.8			
	48 hr	44.3 ± 0.14	0.3	7.7 ± 0.12	1.5	0.026 ± 0.00	4.5			
	Sat.	58.3 ± 0.48	0.8	0.58 ± 0.06	9.5	0.462 ± 0.02	5.7			
<b>Treatment 2</b>	9	10.7 ± 0.21	1.9	0.47 ± 0.03	5.7	0.229 ± 0.03	13.2			
	8	13.7 ± 0.24	1.7	0.43 ± 0.01	2.7	0.171 ± 0.02	13.2			
	7	12.7 ± 0.19	1.5	0.39 ± 0.04	10.7	0.104 ± 0.00	6.6			
	6	14.6 ± 0.06	0.4	0.20 ± 0.03	13.3	0.086 ± 0.00	5.4			
	5	13.2 ± 0.11	0.8	0.40 ± 0.03	7.1	0.116 ± 0.01	14.8			
	48 hr	30.1 ± 0.20	0.7	0.37 ± 0.01	2.9	<0.02				
	Sat.	36.6 ± 0.04	0.1	0.42 ± 0.03	7.9	0.155 ± 0.02	13.3			
<b>Treatment 3</b>	9	12.0 ± 0.11	0.9	0.17 ± 0.01	3.6	0.125 ± 0.01	12.1			
	8	11.5 ± 0.11	1.0	0.23 ± 0.02	9.8	0.076 ± 0.02	28.5			
	7	12.5 ± 0.10	0.8	0.16 ± 0.03	17.6	0.060 ± 0.00	12.9			
	6	12.2 ± 0.11	0.9	0.29 ± 0.03	10.5	0.038 ± 0.00	15.1			
	5	11.1 ± 0.03	0.2	0.21 ± 0.01	5.7	0.040 ± 0.00	10.0			
	48hr	65.3 ± 0.28	0.4	0.06 ± 0.01	14.5	<0.02				
	Sat.	27.6 ± 0.13	0.5	0.13 ± 0.01	9.3	0.055 ± 0.00	7.0			
		Cesium			Barium			Lanthanum		
<b>Treatment 1</b>	9	0.214 ± 0.00	2.2	6.55 ± 0.11	1.7	0.84 ± 0.06	7.7			
	8	0.190 ± 0.00	2.3	6.41 ± 0.09	1.4	<0.1				
	7	0.249 ± 0.00	0.6	5.84 ± 0.12	2.1	0.21 ± 0.01	3.5			
	6	0.197 ± 0.00	1.9	7.09 ± 0.11	1.5	0.34 ± 0.03	8.7			
	5	0.194 ± 0.01	5.0	6.90 ± 0.13	2.0	<0.1				
	48 hr	0.210 ± 0.00	0.9	12.8 ± 0.20	1.6	<0.1				
	Sat.	0.194 ± 0.00	2.0	26.4 ± 0.37	1.4	0.21 ± 0.05	24.2			
<b>Treatment 2</b>	9	0.514 ± 0.00	1.3	5.69 ± 0.06	1.1	<0.1				
	8	0.200 ± 0.00	1.0	6.78 ± 0.03	0.5	0.14 ± 0.04	26.5			
	7	0.186 ± 0.00	1.2	6.15 ± 0.05	0.8	0.28 ± 0.03	9.4			
	6	0.187 ± 0.00	0.2	7.01 ± 0.10	1.4	<0.1				
	5	0.186 ± 0.00	0.8	6.82 ± 0.05	0.8	<0.1				
	48 hr	0.186 ± 0.00	1.7	8.81 ± 0.07	0.8	0.21 ± 0.02	8.7			
	Sat.	0.194 ± 0.00	2.7	18.3 ± 0.14	0.7	0.28 ± 0.02	8.5			
<b>Treatment 3</b>	9	0.199 ± 0.00	2.9	6.73 ± 0.09	1.3	<0.1				
	8	0.194 ± 0.00	2.1	6.25 ± 0.09	1.4	<0.1				
	7	0.204 ± 0.00	0.6	6.84 ± 0.09	1.3	<0.1				
	6	0.189 ± 0.00	1.0	7.08 ± 0.08	1.1	<0.1				
	5	0.202 ± 0.00	0.2	6.01 ± 0.03	0.5	0.19 ± 0.02	10.4			
	48hr	0.194 ± 0.00	2.2	20.3 ± 0.20	1.0	<0.1				
	Sat.	0.199 ± 0.00	1.6	15.7 ± 0.10	0.6	<0.1				

**Table D-5b. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Bone Char (Page 3 of 4)**

of added water	pH	Cerium			Praeseodymium			Samarium			
	ppb	ppb	StD%	ppb	ppb	StD%	ppb	ppb	StD%	StD%	
<b>Treatment 1</b>	9	1.7	± 0.07	4	0.21	± 0.01	5	0.25	± 0.13	51	
	8	0.12	± 0.03	23.9	<0.05			<0.05			
	7	0.4	± 0.02	6.2	<0.05			0.14	± 0.09	65	
	6	0.91	± 0.02	2.6	0.09	± 0.02	20.9	0.1	± 0.06	56	
	5	<0.1			<0.05			0.1	± 0.04	41	
	48 hr	<0.1			<0.05			0.14	± 0.09	66	
	Sat.	0.51	± 0.06	12.5	<0.05			0.16	± 0.05	28	
	<b>Treatment 2</b>	9	0.1	± 0.03	30.1	<0.05		0.11	± 0.09	83	
		8	0.23	± 0.02	9.1	<0.05		0.11	± 0.08	68	
		7	0.31	± 0.05	15.6	0.05	± 0.02	41.5	<0.05		
		6	0.11	± 0.02	15.4	<0.05			<0.05		
		5	0.13	± 0.01	7.1	<0.05			<0.05		
		48 hr	0.34	± 0.06	17.5	<0.05			<0.05		
<b>Treatment 3</b>	Sat.	0.44	± 0.03	7	0.08	± 0.03	37.6	0.1	± 0.06	64	
	9	<0.1			<0.05				<0.05		
	8	0.18	± 0.1	54.4	<0.05			0.07	± 0.02	28	
	7	<0.1			<0.05				<0.05		
	6	0.13	± 0.01	6	<0.05			0.1	± 0.03	31	
	5	0.24	± 0.01	5.4	0.05	± 0.02	27.4	<0.05			
	48 hr	<0.1			<0.05				<0.05		
	Sat.	0.11	± 0.01	13.5	<0.05				<0.05		
	<b>Europium</b>			<b>Gadolinium</b>			<b>Dysprosium</b>				
	<b>Treatment 1</b>	9	0.064	± 0.02	23	0.61	± 0.05	8	0.13	± 0.05	38
		8	<0.05			0.62	± 0.13	21	<0.05		
		7	<0.05			2.29	± 0.26	11	0.15	± 0.06	38
		6	<0.05			0.48	± 0.09	18	0.06	± 0.04	65
		5	<0.05			0.95	± 0.12	13	<0.05		
		48 hr	0.058	± 0.03	55	0.21	± 0.05	25	0.06	± 0.04	67
<b>Treatment 2</b>	Sat.	0.09	± 0	4	0.38	± 0.09	22	<0.05			
	9	<0.05			0.37	± 0.07	20	0.07	± 0.03	50	
	8	<0.05			0.46	± 0.11	23	<0.05			
	7	<0.05			0.43	± 0.08	20	0.1	± 0.03	26	
	6	<0.05			0.23	± 0.02	11	<0.05			
	5	0.092	± 0.01	8	0.44	± 0.05	10	0.11	± 0.11	99	
	48 hr	<0.05			0.16	± 0.09	58	0.19	± 0.07	38	
	Sat.	0.085	± 0.01	16	0.36	± 0.01	4	0.41	± 0.01	3	
	<b>Treatment 3</b>	9	<0.05			0.15	± 0.05	33	0.05	± 0.04	75
		8	<0.05			0.31	± 0.07	22	0.17	± 0.06	38
		7	0.052	± 0.01	9	0.28	± 0.13	49	<0.05		
		6	<0.05			0.23	± 0.05	21	0.14	± 0.06	45
		5	0.053	± 0.03	56	0.19	± 0.01	7	0.31	± 0.03	8
		48 hr	0.056	± 0.04	68	0.05	± 0.02	38	0.08	± 0.09	111
SSat.			<0.05			0.1	± 0.01	12	0.08	± 0.04	44

**Table D-5b. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Bone Char (Page 4 of 4)**

	<b>pH of added water</b>	<b>Holmium</b>			<b>Erbium</b>			<b>Lead</b>			<b>StD%</b>		
		<b>ppb</b>	<b>ppb</b>	<b>StD%</b>	<b>ppb</b>	<b>ppb</b>	<b>StD%</b>	<b>ppb</b>	<b>ppb</b>	<b>StD%</b>			
<b>Treatment 1</b>	9	0.067	±	0.02	31	0.16	±	0.06	34	0.74	±	0.01	1.3
	8	<0.02				<0.05				0.39	±	0.03	7.2
	7	0.031	±	0.02	66	0.06	±	0.02	30	0.58	±	0.02	4
	6	<0.02				0.06	±	0.02	36	0.37	±	0.01	3
	5	<0.02				0.06	±	0.06	92	0.27	±	0.02	5.9
	48 hr	0.021	±	0.02	79	0.07	±	0.01	16	0.16	±	0.02	11.6
	Sat.	<0.02				0.09	±	0.02	17	0.43	±	0.01	3.3
<b>Treatment 2</b>	9	<0.02				<0.05				0.36	±	0.01	2.4
	8	<0.02				<0.05				0.41	±	0.03	6
	7	<0.02				<0.05				0.33	±	0.02	5.5
	6	<0.02				<0.05				0.19	±	0.02	7.7
	5	<0.02				<0.05				0.4	±	0.03	6.1
	48 hr	0.033	±	0.03	92	<0.05				0.45	±	0.04	9.4
	Sat.	<0.02				0.06	±	0.03	61	0.89	±	0.02	1.9
<b>Treatment 3</b>	9	<0.02				<0.05				0.12	±	0.01	11
	8	0.025	±	0.01	24	<0.05				0.43	±	0.02	5.6
	7	<0.02				<0.05				0.12	±	0.01	6.3
	6	0.023	±	0.01	30	0.05	±	0.03	50	0.3	±	0	0.6
	5	0.023	±	0.02	69	<0.05				0.46	±	0.02	3.7
	48 hr	<0.02				<0.05				0.11	±	0.02	15.1
	Sat.	0.04	±	0.02	40	0.06	±	0.02	33	0.16	±	0.02	9.2

**Table D-5c. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Hydroxyapatite (Page 1 of 4)**

	pH	Chromium			Manganese			Iron		
	of added water	ppb	ppb	StD%	ppb	ppb	StD%	ppb	ppb	StD%
<b>Treatment 1</b>	9	10.1 ± 0.16		1.6	1.68 ± 0.02	0.02	1.4	467 ± 7.5	7.5	1.6
	8	13.8 ± 0.36		2.6	1.46 ± 0.02	0.02	1.6	407 ± 20.4	20.4	5.0
	7	14.1 ± 0.31		2.2	1.60 ± 0.01	0.01	0.8	398 ± 5.3	5.3	1.3
	6	15.0 ± 0.12		0.8	1.61 ± 0.05	0.05	3.1	468 ± 8.3	8.3	1.8
	5	13.5 ± 0.13		1.0	1.46 ± 0.02	0.02	1.1	494 ± 7.8	7.8	1.6
	48 hr	0.3 ± 0.02		8.5	2.06 ± 0.03	0.03	1.7	349 ± 2.9	2.9	0.8
	Satur	35.8			2.39			669		
<b>Treatment 2</b>	9	18.8 ± 0.05		0.2	1.19 ± 0.01	0.01	0.9	403 ± 4.1	4.1	1.0
	8	20.2 ± 0.10		0.5	1.76 ± 0.04	0.04	2.5	462 ± 8.2	8.2	1.8
	7	19.0 ± 0.32		1.7	1.62 ± 0.01	0.01	0.3	463 ± 3.9	3.9	0.8
	6	18.4 ± 0.28		1.5	1.73 ± 0.03	0.03	1.6	468 ± 10.5	10.5	2.3
	5	18.3 ± 0.19		1.0	1.51 ± 0.03	0.03	1.9	467 ± 4.7	4.7	1.0
	48 hr	7.4 ± 0.12		1.7	1.87 ± 0.01	0.01	0.6	299 ± 2.2	2.2	0.7
	Satur	38.2			3.62			614		
<b>Treatment 3</b>	9	7.8 ± 0.09		1.2	1.39 ± 0.02	0.02	1.4	341 ± 5.6	5.6	1.7
	8	7.3 ± 0.11		1.5	1.05 ± 0.00	0.00	0.1	350 ± 6.4	6.4	1.8
	7	7.0 ± 0.10		1.4	1.24 ± 0.01	0.01	1.1	386 ± 6.5	6.5	1.7
	6	6.8 ± 0.09		1.3	1.32 ± 0.01	0.01	0.5	391 ± 12.9	12.9	3.3
	5	6.7 ± 0.03		0.4	1.45 ± 0.02	0.02	1.6	425 ± 1.8	1.8	0.4
	48 hr	1.9 ± 0.01		0.3	1.12 ± 0.01	0.01	1.2	291 ± 0.7	0.7	0.2
	Satur	10.4 ± 0.05		0.4	1.42 ± 0.01	0.01	1.0	348 ± 5.4	5.4	1.5
Copper      Zinc      Arsenic										
<b>Treatment 1</b>	9	1.39 ± 0.03		2.5	9.85 ± 0.30	0.30	3.0	0.11 ± 0.00	0.00	2.8
	8	0.89 ± 0.02		2.1	3.67 ± 0.13	0.13	3.5	0.15 ± 0.01	0.01	6.7
	7	2.00 ± 0.01		0.4	2.98 ± 0.09	0.09	2.9	0.13 ± 0.00	0.00	3.6
	6	1.24 ± 0.10		8.0	19.2 ± 0.47	0.47	2.4	0.14 ± 0.01	0.01	8.4
	5	0.98 ± 0.06		6.1	4.49 ± 0.05	0.05	1.2	0.14 ± 0.00	0.00	1.9
	48 hr	0.47 ± 0.05		10.4	1.74 ± 0.15	0.15	8.7	0.12 ± 0.02	0.02	13.4
	Satur	4.34			10.3			0.43		
<b>Treatment 2</b>	9	1.83 ± 0.04		2.3	2.73 ± 0.08	0.08	2.8	0.23 ± 0.01	0.01	3.5
	8	2.01 ± 0.04		2.1	8.35 ± 0.24	0.24	2.8	0.22 ± 0.02	0.02	7.9
	7	2.34 ± 0.02		0.9	2.03 ± 0.06	0.06	3.0	0.24 ± 0.02	0.02	6.9
	6	1.09 ± 0.06		5.7	6.40 ± 0.21	0.21	3.2	0.21 ± 0.01	0.01	2.6
	5	1.63 ± 0.07		4.2	2.82 ± 0.07	0.07	2.3	0.21 ± 0.04	0.04	17.2
	48 hr	0.94 ± 0.04		4.6	6.04 ± 0.04	0.04	0.7	0.16 ± 0.03	0.03	17.7
	Satur	5.50			64.0 ± 0.00	0.00	0.0	0.59 ± 0.00	0.00	0.0
<b>Treatment 3</b>	9	1.84 ± 0.04		2.4	5.94 ± 0.06	0.06	1.0	0.26 ± 0.03	0.03	12.2
	8	1.33 ± 0.09		7.1	6.58 ± 0.04	0.04	0.6	0.24 ± 0.03	0.03	14.0
	7	1.13 ± 0.02		1.5	1.57 ± 0.11	0.11	7.3	0.22 ± 0.03	0.03	14.1
	6	2.12 ± 0.05		2.1	2.98 ± 0.15	0.15	4.9	0.19 ± 0.02	0.02	9.8
	5	1.09 ± 0.02		1.8	1.78 ± 0.05	0.05	3.0	0.21 ± 0.02	0.02	8.3
	48 hr	1.31 ± 0.08		6.3	8.05 ± 0.10	0.10	1.2	0.14 ± 0.03	0.03	21.0
	Satur	3.15 ± 0.13		4.0	8.28 ± 0.17	0.17	2.0	0.43 ± 0.02	0.02	4.0

**Table D-5c. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Hydroxyapatite (Page 2 of 4)**

of added water	pH	Strontium		Yttrium			Silver	
	ppb	StD%	ppb	StD%	ppb	StD%	ppb	StD%
<b>Treatment 1</b>	9	88.3 ± 0.54	0.6	<0.05			<0.02	
	8	83.9 ± 0.64	0.8	<0.05			<0.02	
	7	76.3 ± 0.38	0.5	<0.05			<0.02	
	6	90.8 ± 0.65	0.7	0.07 ± 0.02	36.7	<0.02		
	5	93.7 ± 0.28	0.3	0.07 ± 0.01	10.9	<0.02		
	48 hr	55.9 ± 0.50	0.9	<0.05			<0.02	
	Satur	118		not			<0.02	
<b>Treatment 2</b>	9	81.1 ± 0.83	1.0	0.17 ± 0.02	14.8	<0.02		
	8	90.6 ± 0.70	0.8	0.35 ± 0.01	2.6	<0.02		
	7	92.1 ± 0.01	0.0	not			<0.02	
	6	90.7 ± 0.82	0.9	0.14 ± 0.01	9.5	<0.02		
	5	90.3 ± 0.30	0.3	0.09 ± 0.02	19.4	<0.02		
	48 hr	41.4 ± 0.15	0.4	0.15 ± 0.04	24.2	<0.02		
	Satur	103 ± 0.00	0.0	not			<0.02	
<b>Treatment 3</b>	9	69.4 ± 0.24	0.3	0.14 ± 0.01	8.3	0.028 ± 0.00		23.7
	8	69.6 ± 0.19	0.3	0.17 ± 0.02	12.7	<0.02		
	7	78.7 ± 0.41	0.5	0.09 ± 0.01	8.8	<0.02		
	6	76.8 ± 0.30	0.4	0.07 ± 0.02	23.7	<0.02		
	5	82.6 ± 0.42	0.5	0.07 ± 0.01	10.6	0.038 ± 0.00		14.5
	48 hr	49.4 ± 0.18	0.4	0.10 ± 0.02	24.8	<0.02		
	Satur	62.0 ± 0.19	0.3	0.09 ± 0.01	11.1	0.042 ± 0.00		20.1
<b>Cesium</b>								
<b>Treatment 1</b>	9	0.189 ± 0.00	0.4	22.2 ± 0.02	0.1	<0.1		
	8	0.202 ± 0.00	1.5	23.8 ± 0.19	0.8	<0.1		
	7	0.222 ± 0.00	0.4	20.9 ± 0.14	0.7	<0.1		
	6	0.193 ± 0.00	2.8	27.2 ± 0.29	1.1	<0.1		
	5	0.193 ± 0.00	1.5	27.7 ± 0.13	0.5	<0.1		
	48 hr	0.193 ± 0.00	0.5	13.1 ± 0.13	1.0	<0.1		
	Satur	0.201 ± 0.00	0.0	31.3		0.11		
<b>Treatment 2</b>	9	0.192 ± 0.00	0.6	21.9 ± 0.10	0.5	<0.1		
	8	0.196 ± 0.00	1.5	24.9 ± 0.17	0.7	<0.1		
	7	0.194 ± 0.00	0.8	25.4 ± 0.15	0.6	<0.1		
	6	0.190 ± 0.00	0.9	25.0 ± 0.11	0.4	<0.1		
	5	0.199 ± 0.00	1.2	29.2 ± 0.08	0.3	<0.1		
	48hr	0.194 ± 0.00	0.2	9.72 ± 0.02	0.2	<0.1		
	Satur	0.195 ± 0.00	0.0	29.8		0.21		
<b>Treatment 3</b>	9	0.194 ± 0.00	0.6	20.8 ± 0.04	0.2	<0.1		
	8	0.192 ± 0.00	1.6	19.6 ± 0.25	1.3	<0.1		
	7	0.213 ± 0.00	1.8	22.7 ± 0.29	1.3	<0.1		
	6	0.197 ± 0.00	1.3	22.3 ± 0.15	0.7	<0.1		
	5	0.203 ± 0.00	1.3	24.3 ± 0.10	0.4	<0.1		
	48 hr	0.235 ± 0.00	1.0	12.4 ± 0.06	0.4	<0.1		
	Satur	0.200 ± 0.00	1.0	3.79 ± 0.07	1.8	<0.1		

**Table D-5c. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Hydroxyapatite (Page 3 of 4)**

	pH	Cerium		Praeseodymium		Samarium	
	of added water	ppb	StD%	ppb	StD%	ppb	StD%
<b>Treatment 1</b>	9	<0.1		<0.05		<0.05	
	8	<0.1		<0.05		0.07 ± 0.05	69
	7	<0.1		<0.05		<0.05	
	6	<0.1		<0.05		<0.05	
	5	<0.1		<0.05		0.07 ± 0.04	62
	48 hr	<0.1		<0.05		<0.05	
	Satur	0.23		0.06		<0.05	
	9	<0.1		<0.05		0.08 ± 0.02	30
	8	<0.1		<0.05		0.07 ± 0.05	70
	7	<0.1		<0.05		<0.05	
<b>Treatment 2</b>	6	0.10 ± 0.00	3.5	<0.05		0.07 ± 0.05	72
	5	<0.1		<0.05		<0.05	
	48 hr	0.13 ± 0.02	16.3	<0.05		<0.05	
	Satur	0.30		0.07		0.10	
	9	<0.1		<0.05		0.09 ± 0.03	31
	8	<0.1		<0.05		<0.05	
	7	<0.1		<0.05		<0.05	
<b>Treatment 3</b>	6	<0.1		<0.05		0.06 ± 0.02	43
	5	<0.1		<0.05		0.05 ± 0.01	23
	48 hr	<0.1		<0.05		<0.05	
	Satur	<0.1		<0.05		0.08 ± 0.01	19
	9	<0.05		0.06 ± 0.05	81	<0.05	
	8	<0.05		0.14 ± 0.01	5	<0.05	
	7	0.064 ± 0.02	44	0.04 ± 0.02	49	<0.05	
<b>Treatment 1</b>	6	0.072 ± 0.03	44	<0.02		<0.05	
	5	<0.05		0.03 ± 0.05	155	<0.05	
	48 hr	<0.05		0.06 ± 0.02	44	<0.05	
	Satur	0.077		0.11		<0.05	
	9	0.056 ± 0.01	28	<0.02		<0.05	
	8	0.067 ± 0.00	11	0.07 ± 0.01	14	0.10 ± 0.01	13
	7	0.052 ± 0.00	6	0.09 ± 0.09	103	<0.05	
<b>Treatment 2</b>	6	0.056 ± 0.03	59	0.07 ± 0.03	42	<0.05	
	5	0.056 ± 0.02	46	<0.02		<0.05	
	48 hr	<0.05		0.02 ± 0.02	97	0.10 ± 0.08	78
	Satur	0.112		0.08		0.53	
	9	<0.05		0.02 ± 0.02	115	0.09 ± 0.01	15
	8	<0.05		0.06 ± 0.02	34	0.19 ± 0.07	36
	7	<0.05		0.04 ± 0.03	83	<0.05	
<b>Treatment 3</b>	6	0.069 ± 0.02	38	0.09 ± 0.05	51	<0.05	
	5	0.061 ± 0.00	14	0.06 ± 0.03	52	<0.05	
	48 hr	<0.05		0.05 ± 0.01	29	0.07 ± 0.01	17
	Satur	<0.05		0.16 ± 0.03	19	<0.05	
	9	<0.05		0.02 ± 0.02	115	0.09 ± 0.01	15
	8	<0.05		0.06 ± 0.02	34	0.19 ± 0.07	36
	7	<0.05		0.04 ± 0.03	83	<0.05	

**Table D-5c. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Hydroxyapatite (Page 4 of 4)**

	pH of added water	Holmium		Erbium		Lead	
		ppb	StD%	ppb	StD%	ppb	StD%
<b>Treatment 1</b>	9	<0.02		<0.05		<0.1	
	8	<0.02		<0.05		0.12 ± 0.00	5.0
	7	<0.02		<0.05		0.16 ± 0.01	11.9
	6	<0.02		<0.05		0.13 ± 0.01	13.7
	5	<0.02		<0.05		<0.1	
	48 hr	<0.02		<0.05		<0.1	
	Satur	<0.02		<0.05		<0.1	
<b>Treatment 2</b>	9	<0.02		<0.05		<0.1	
	8	<0.02		<0.05		0.29 ± 0.02	9.0
	7	<0.02		<0.05		<0.1	
	6	<0.02		<0.05		<0.1	
	5	<0.02		<0.05		<0.1	
	48 hr	<0.02		<0.05		0.64 ± 0.05	8.1
	Satur	0.029		<0.05		1.02	
<b>Treatment 3</b>	9	<0.02		<0.05		<0.1	
	8	<0.02		<0.05		0.21 ± 0.01	6.5
	7	<0.02		<0.05		<0.1	
	6	<0.02		<0.05		<0.1	
	5	<0.02		<0.05		<0.1	
	48 hr	0.021 ± 0.01	50	<0.05		0.18 ± 0.01	9.5
	Satur	<0.02		0.06 ± 0.01	19	0.16 ± 0.01	10.2

**Table D-5d. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Clinoptilolite (Page 1 of 4)**

	pH	Chromium		Manganese		Iron		
	of added water	ppb	StD%	ppb	StD%	ppb	StD%	
<b>Treatment 1</b>	9	8.0 ± 0.13	1.6	12.6 ± 0.11	0.9	409 ± 1.2	0.3	
	8	5.0 ± 0.08	1.6	5.18 ± 0.01	0.3	193 ± 1.6	0.8	
	7	4.7 ± 0.08	1.8	12.5 ± 0.21	1.6	429 ± 6.4	1.5	
	6	4.5 ± 0.05	1.0	4.88 ± 0.05	1.0	183 ± 0.8	0.4	
	5	4.9 ± 0.02	0.4	4.56 ± 0.01	0.2	226 ± 1.2	0.5	
	48 hr	0.9 ± 0.03	3.2	34.1 ± 0.17	0.5	923 ± 7.3	0.8	
	Sat.	22.2 ± 0.14	0.6	16.7 ± 0.16	1.0	557 ± 6.1	1.1	
	<b>Treatment 2</b>	9	10.8 ± 0.07	0.6	3.86 ± 0.06	1.7	145 ± 4.3	3.0
	8	9.7 ± 0.03	0.3	1.93 ± 0.04	2.2	92 ± 1.6	1.7	
	7	9.8 ± 0.07	0.7	2.52 ± 0.04	1.5	110 ± 1.5	1.3	
<b>Treatment 3</b>	6	10.1 ± 0.10	1.0	3.03 ± 0.04	1.4	119 ± 0.9	0.8	
	5	10.1 ± 0.07	0.7	3.41 ± 0.03	0.8	130 ± 0.8	0.6	
	48 hr	2.2 ± 0.04	1.9	55.6 ± 0.14	0.2	1406 ± 2.5	0.2	
	Sat.	50.9 ± 0.31	0.6	20.1 ± 0.11	0.6	590 ± 5.9	1.0	
	9	1.5 ± 0.03	1.9	3.35 ± 0.02	0.7	124 ± 1.6	1.3	
	8	2.9 ± 0.10	3.4	2.89 ± 0.02	0.8	106 ± 0.5	0.5	
	7	2.0 ± 0.02	1.2	9.17 ± 0.02	0.2	195 ± 3.4	1.8	
	6	1.8 ± 0.03	1.7	2.72 ± 0.05	1.9	90 ± 0.7	0.8	
	5	1.4 ± 0.14	9.8	3.38 ± 0.38	11.1	168 ± 25.9	15.4	
	48 hr	3.1 ± 0.03	0.8	28.1 ± 0.07	0.3	754 ± 13.0	1.7	
	Sat.	14.0 ± 0.03	0.2	33.1 ± 0.02	0.1	796 ± 4.8	0.6	
		Copper		Zin		Arsenic		
<b>Treatment 1</b>	9	15.2 ± 0.24	1.6	32.4 ± 0.32	1.0	21.7 ± 0.49	2.3	
	8	6.12 ± 0.08	1.3	3.79 ± 0.10	2.6	11.3 ± 0.07	0.6	
	7	7.53 ± 0.10	1.3	3.49 ± 0.03	0.9	9.99 ± 0.34	3.4	
	6	5.84 ± 0.03	0.6	4.05 ± 0.05	1.2	12.7 ± 0.23	1.8	
	5	5.69 ± 0.06	1.1	5.70 ± 0.10	1.7	11.9 ± 0.33	2.8	
	48 hr	3.80 ± 0.03	0.7	12.6 ± 0.15	1.2	1.25 ± 0.09	7.3	
	Sat.	14.8 ± 0.21	1.4	42.5 ± 1.47	3.4	20.5 ± 0.64	3.1	
	<b>Treatment 2</b>	9	7.58 ± 0.12	1.6	15.1 ± 0.05	0.3	14.7 ± 0.18	1.2
	8	4.54 ± 0.06	1.2	1.98 ± 0.09	4.5	10.2 ± 0.19	1.8	
	7	5.64 ± 0.07	1.2	1.69 ± 0.08	4.8	12.4 ± 0.35	2.9	
<b>Treatment 3</b>	6	4.48 ± 0.06	1.4	7.59 ± 0.20	2.7	10.6 ± 0.13	1.2	
	5	5.05 ± 0.04	0.9	4.11 ± 0.16	4.0	12.4 ± 0.37	3.0	
	48 hr	5.77 ± 0.12	2.1	14.1 ± 0.19	1.3	1.47 ± 0.14	9.8	
	Sat.	14.6 ± 0.10	0.7	16.5 ± 0.12	0.7	37.5 ± 0.46	1.2	
	9	8.11 ± 0.12	1.5	2.85 ± 0.23	8.0	11.5 ± 0.25	2.2	
	8	5.37 ± 0.04	0.7	7.51 ± 0.30	4.0	8.44 ± 0.19	2.3	
	7	5.54 ± 0.16	2.9	5.68 ± 0.12	2.0	9.77 ± 0.18	1.9	
	6	4.72 ± 0.10	2.1	6.81 ± 0.19	2.8	9.76 ± 0.19	1.9	
	5	5.08 ± 0.61	12.0	7.69 ± 1.30	16.9	9.07 ± 1.23	13.5	
	48 hr	4.02 ± 0.01	0.3	8.04 ± 0.11	1.4	1.68 ± 0.05	2.7	
	Sat.	15.1 ± 0.12	0.8	13.0 ± 0.26	2.0	33.6 ± 0.53	1.6	

**Table D-5d. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Clinoptilolite (Page 2 of 4)**

of added water	pH	Strontium		Yttrium		Silver		StD%
	ppb	ppb	StD%	ppb	StD%	ppb	ppb	
<b>Treatment 1</b>	9	4.6 ± 0.09	1.9	43.8 ± 0.86	2.0	0.050 ± 0.00	0.00	4.8
	8	2.6 ± 0.06	2.2	28.2 ± 0.49	1.7	0.052 ± 0.00	0.00	6.3
	7	3.0 ± 0.01	0.3	24.3 ± 0.26	1.1	0.089 ± 0.01	0.01	19.2
	6	2.9 ± 0.02	0.6	58.6 ± 0.39	0.7	0.070 ± 0.00	0.00	9.7
	5	2.7 ± 0.01	0.5	20.5 ± 0.48	2.4	0.068 ± 0.00	0.00	13.4
	48 hr	3.8 ± 0.08	2.0	2.17 ± 0.11	4.9	0.029 ± 0.00	0.00	11.5
	Sat.	16.8 ± 0.21	1.2	28.2 ± 0.17	0.6	0.055 ± 0.00	0.00	3.9
<b>Treatment 2</b>	9	3.0 ± 0.04	1.2	1.88 ± 0.06	3.3	0.172 ± 0.01	0.01	9.7
	8	2.5 ± 0.02	0.8	0.58 ± 0.08	13.9	0.047 ± 0.00	0.00	7.6
	7	2.9 ± 0.02	0.6	0.61 ± 0.02	3.1	0.113 ± 0.01	0.01	11.5
	6	2.6 ± 0.01	0.5	0.70 ± 0.04	5.7	0.078 ± 0.00	0.00	5.2
	5	3.7 ± 0.06	1.6	1.87 ± 0.08	4.4	0.108 ± 0.01	0.01	10.1
	48 hr	3.7 ± 0.04	1.2	4.06 ± 0.14	3.5	<0.02		
	Sat.	11.3 ± 0.08	0.7	2.33 ± 0.02	0.9	0.108 ± 0.00	0.00	8.1
<b>Treatment 3</b>	9	2.5 ± 0.01	0.6	1.00 ± 0.03	3.0	0.249 ± 0.01	0.01	5.9
	8	2.4 ± 0.03	1.1	0.71 ± 0.02	3.3	0.116 ± 0.00	0.00	5.6
	7	3.2 ± 0.03	0.9	1.72 ± 0.10	5.6	0.079 ± 0.00	0.00	3.1
	6	2.2 ± 0.04	1.7	0.61 ± 0.04	5.8	0.187 ± 0.02	0.02	12.1
	5	2.9 ± 0.50	17.2	0.86 ± 0.08	9.6	0.158 ± 0.03	0.03	20.6
	48hr	3.6 ± 0.05	1.4	2.64 ± 0.07	2.8	0.054 ± 0.00	0.00	17.1
	Sat.	11.5 ± 0.19	1.7	4.30 ± 0.09	2.2	0.145 ± 0.00	0.00	3.2
<b>Cesium</b>								
<b>Treatment 1</b>	9	0.237 ± 0.00	2.7	3.16 ± 0.12	3.8	2.40 ± 0.06	0.06	2.5
	8	0.230 ± 0.00	1.8	1.52 ± 0.03	1.8	1.48 ± 0.04	0.04	2.4
	7	0.242 ± 0.00	0.4	3.04 ± 0.09	3.0	2.77 ± 0.44	0.44	15.8
	6	0.230 ± 0.00	1.3	1.57 ± 0.03	2.2	1.28 ± 0.10	0.10	7.7
	5	0.241 ± 0.00	1.1	1.40 ± 0.04	2.7	1.20 ± 0.05	0.05	3.9
	48 hr	0.230 ± 0.00	0.3	6.05 ± 0.05	0.8	3.17 ± 0.16	0.16	5.1
	Sat.	0.316 ± 0.00	1.5	5.19 ± 0.01	0.3	2.39 ± 0.08	0.08	3.1
<b>Treatment 2</b>	9	0.317 ± 0.00	1.7	1.17 ± 0.01	1.0	1.05 ± 0.04	0.04	4.1
	8	0.233 ± 0.00	0.7	0.75 ± 0.01	1.1	0.53 ± 0.05	0.05	10.1
	7	0.237 ± 0.00	2.3	0.93 ± 0.04	4.1	0.66 ± 0.08	0.08	12.1
	6	0.237 ± 0.00	1.4	1.03 ± 0.01	1.4	0.74 ± 0.02	0.02	2.5
	5	0.246 ± 0.00	1.2	3.30 ± 0.04	1.3	0.94 ± 0.14	0.14	15.1
	48 hr	0.234 ± 0.00	1.6	8.64 ± 0.09	1.0	5.56 ± 0.12	0.12	2.1
	Sat.	0.246 ± 0.00	1.1	7.81 ± 0.10	1.2	2.61 ± 0.04	0.04	1.5
<b>Treatment 3</b>	9	0.259 ± 0.00	1.5	3.23 ± 0.00	0.0	1.04 ± 0.09	0.09	8.6
	8	0.240 ± 0.00	3.1	0.94 ± 0.02	2.3	0.88 ± 0.09	0.09	9.7
	7	0.326 ± 0.00	2.1	1.83 ± 0.07	4.0	2.00 ± 0.05	0.05	2.4
	6	0.242 ± 0.00	1.7	0.80 ± 0.03	3.3	0.82 ± 0.05	0.05	6.5
	5	0.534 ± 0.14	27.4	1.75 ± 0.23	13.3	0.87 ± 0.12	0.12	13.5
	48 hr	0.232 ± 0.00	1.4	6.11 ± 0.05	0.7	3.32 ± 0.23	0.23	7.0
	Sat.	0.273 ± 0.00	1.1	7.80 ± 0.02	0.3	4.39 ± 0.18	0.18	4.1

**Table D-5d. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Clinoptilolite (Page 3 of 4)**

	pH of added water	Cerium		Praeseodymium			Samarium			StD%	
		ppb	StD%	ppb	StD%	ppb	StD%	ppb	StD%		
<b>Treatment 1</b>	9	5.34	± 0.21	4.0	0.57	± 0.08	15.5	0.43	± 0.15	36	
	8	2.92	± 0.07	2.5	0.31	± 0.04	15.8	0.36	± 0.15	42	
	7	5.48	± 0.04	0.7	0.60	± 0.02	3.4	0.55	± 0.12	21	
	6	2.82	± 0.07	2.6	0.30	± 0.01	4.6	0.21	± 0.09	42	
	5	2.87	± 0.12	4.4	0.26	± 0.02	10.7	0.28	± 0.07	25	
	48 hr	11.95	± 0.28	2.3	0.78	± 0.00	0.7	0.81	± 0.24	30	
	Sat.	6.44	± 0.17	2.7	0.64	± 0.02	4.3	0.81	± 0.32	39	
	<b>Treatment 2</b>	9	2.51	± 0.08	3.3	0.22	± 0.01	6.3	0.43	± 0.03	8
		8	1.33	± 0.13	9.5	0.13	± 0.01	13.1	0.23	± 0.07	31
		7	1.72	± 0.05	3.2	0.17	± 0.01	7.5	0.23	± 0.19	85
		6	1.94	± 0.06	3.2	0.17	± 0.01	9.5	0.22	± 0.07	33
		5	2.13	± 0.05	2.4	0.24	± 0.00	2.2	0.41	± 0.27	65
		48 hr	20.20	± 0.48	2.4	1.40	± 0.05	3.7	1.36	± 0.07	5
<b>Treatment 3</b>	Sat.	7.96	± 0.06	0.7	0.72	± 0.05	7.7	1.03	± 0.18	17	
	9	2.21	± 0.07	3.2	0.24	± 0.03	12.8	0.11	± 0.11	95	
	8	1.91	± 0.11	5.8	0.20	± 0.01	6.5	0.13	± 0.12	92	
	7	4.94	± 0.16	3.2	0.51	± 0.02	4.2	0.42	± 0.14	35	
	6	2.08	± 0.15	7.2	0.19	± 0.03	19.2	0.20	± 0.07	33	
	5	2.52	± 0.64	25.2	0.25	± 0.01	5.0	0.41	± 0.23	56	
	48 hr	11.33	± 0.19	1.7	0.88	± 0.09	10.6	0.78	± 0.10	13	
	Sat.	13.23	± 0.39	2.9	1.19	± 0.03	2.8	1.23	± 0.19	15	
	<b>Europium</b>			<b>Gadolinium</b>			<b>Dysprosium</b>				
	<b>Treatment 1</b>	9	0.154	± 0.03	24	3.74	± 0.18	5	0.73	± 0.11	14
		8	<0.05			1.20	± 0.10	9	0.35	± 0.12	33
		7	0.096	± 0.03	34	0.81	± 0.04	5	0.63	± 0.08	12
		6	0.060	± 0.01	21	6.19	± 0.22	4	0.24	± 0.06	26
		5	<0.05			0.98	± 0.07	7	0.33	± 0.03	9
		48 hr	0.112	± 0.03	31	0.89	± 0.07	7	0.75	± 0.11	15
<b>Treatment 2</b>	Sat.	0.124	± 0.03	25	3.91	± 0.07	2	0.64	± 0.11	17	
	9	0.057	± 0.02	50	3.35	± 0.28	8	1.11	± 0.07	6	
	8	<0.05			0.45	± 0.06	12	0.65	± 0.03	5	
	7	<0.05			1.12	± 0.14	12	0.81	± 0.10	13	
	6	<0.05			0.56	± 0.13	23	1.04	± 0.09	8	
	5	0.070	± 0.04	62	5.17	± 0.15	3	0.75	± 0.08	11	
	48 hr	0.214	± 0.02	11	1.55	± 0.14	9	1.42	± 0.19	13	
	Sat.	0.181	± 0.03	22	1.73	± 0.13	7	2.10	± 0.10	5	
	<b>Treatment 3</b>	9	<0.05			2.91	± 0.16	6	0.54	± 0.01	2
		8	<0.05			0.58	± 0.02	3	0.68	± 0.04	6
		7	0.065	± 0.03	49	1.60	± 0.10	6	0.65	± 0.03	5
		6	<0.05			0.31	± 0.06	19	0.59	± 0.05	9
		5	0.069	± 0.01	27	1.34	± 0.24	18	0.76	± 0.05	7
		48 hr	0.138	± 0.01	12	0.76	± 0.09	12	0.70	± 0.12	17
	Sat.	0.203	± 0.00	4	2.64	± 0.26	10	1.35	± 0.05	4	

**Table D-5d. Inductively Coupled Plasma-Mass Spectrometry Data for the  
pH Stability Test: Clinoptilolite (Page 4 of 4)**

	pH of added water	Holmium			Erbium			Lead			StD%
		ppb	StD%	ppb	StD%	ppb	StD%	ppb	StD%	ppb	
<b>Treatment 1</b>	9	0.099 ± 0.017	17	0.40 ± 0.03	8	1.18 ± 0.05	4.3				
	8	0.034 ± 0.008	23	0.16 ± 0.05	31	0.31 ± 0.02	8.5				
	7	0.094 ± 0.024	26	0.40 ± 0.02	5	0.42 ± 0.02	5.1				
	6	0.050 ± 0.011	22	0.16 ± 0.05	34	0.43 ± 0.00	0.8				
	5	0.056 ± 0.016	29	0.11 ± 0.10	89	0.35 ± 0.02	5.8				
	48 hr	0.110 ± 0.004	3	0.49 ± 0.08	15	0.62 ± 0.00	1.4				
	Sat.	0.102 ± 0.021	20	0.37 ± 0.08	21	1.05 ± 0.01	1.2				
<b>Treatment 2</b>	9	0.044 ± 0.010	22	0.13 ± 0.03	27	0.23 ± 0.01	5.7				
	8	<0.02		0.05 ± 0.01	13	0.22 ± 0.00	2.1				
	7	0.024 ± 0.012	48	0.04 ± 0.03	83	0.16 ± 0.00	2.5				
	6	0.032 ± 0.009	28	0.09 ± 0.05	54	0.23 ± 0.01	7.3				
	5	0.053 ± 0.036	67	0.15 ± 0.10	68	0.29 ± 0.00	3.1				
	48 hr	0.200 ± 0.016	8	0.79 ± 0.11	14	1.55 ± 0.05	3.4				
	Sat.	0.153 ± 0.011	7	0.41 ± 0.09	23	0.80 ± 0.05	7.1				
<b>Treatment 3</b>	9	<0.02		0.10 ± 0.05	53	0.21 ± 0.02	9.5				
	8	0.034 ± 0.012	36	0.16 ± 0.04	24	0.43 ± 0.02	5.3				
	7	0.080 ± 0.014	17	0.20 ± 0.04	18	0.41 ± 0.04	10.1				
	6	0.043 ± 0.012	27	0.06 ± 0.04	63	0.28 ± 0.01	4.3				
	5	0.055 ± 0.043	78	0.19 ± 0.02	12	0.27 ± 0.01	5.8				
	48 hr	0.123 ± 0.032	26	0.32 ± 0.12	36	0.68 ± 0.03	5.5				
	Sat.	0.176 ± 0.021	12	0.55 ± 0.05	9	2.68 ± 0.02	0.8				

**Table D-5e. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Hanford Soil (Page 1 of 4)**

	pH	Chromium			Manganese			Iron					
	of added water	ppb	ppb	StD%	ppb	ppb	StD%	ppb	ppb	StD%			
<b>Treatment 1</b>	9	6.1	±	0.05	0.8	2.53	±	0.04	1.5	232	±	3.0	1.3
	8	5.5	±	0.07	1.3	1.62	±	0.02	1.5	207	±	5.7	2.8
	7	5.8	±	0.27	4.7	5.17	±	0.18	3.5	242	±	11.0	4.6
	6	0.3	±	0.02	6.9	0.44	±	0.01	3.4	172	±	1.9	1.1
	5	0.3	±	0.03	9.1	0.22	±	0.01	5.2	108	±	3.1	2.9
	48 hr	0.4	±	0.02	4.2	8.66	±	0.09	1.1	206	±	8.5	4.1
<b>Treatment 2</b>	9	92.1	±	0.87	0.9	1.09	±	0.01	1.2	243	±	2.3	1.0
	8	88.6	±	0.76	0.9	0.79	±	0.01	1.7	234	±	11.2	4.8
	7	95.2				0.64				245			
	6	102.4	±	0.6	0.6	0.86	±	0.02	2.5	284	±	3.0	1.1
	5	111.9				0.75				286			
	48 hr	3.1	±	0.10	3.2	5.10	±	0.07	1.4	171	±	6.1	3.6
<b>Treatment 3</b>	9	4.7	±	0.08	1.6	1.63	±	0.03	1.8	179	±	6.4	3.6
	8	3.5	±	0.05	1.4	1.52	±	0.01	1.0	205	±	1.8	0.9
	7	3.9	±	0.03	0.7	2.34	±	0.03	1.4	222	±	3.5	1.6
	6	4.1				1.82				220			
	5	3.2				2.08				256			
	48 hr	1.2	±	0.03	2.0	2.21	±	0.02	1.0	198	±	4.2	2.1
		Copper			Zinc			Arsenic					
<b>Treatment 1</b>	9	6.35	±	0.08	1.2	4.78	±	0.14	3.0	1.49	±	0.02	1.3
	8	6.08	±	0.16	2.6	6.97	±	0.46	6.6	1.47	±	0.04	2.8
	7	4.81	±	0.07	1.4	8.80	±	0.22	2.5	1.74	±	0.00	0.3
	6	3.53	±	0.08	2.3	5.14	±	0.05	0.9	<0.05			
	5	1.30	±	0.08	6.3	3.12	±	0.08	2.7	<0.05			
	48 hr	2.88	±	0.08	2.7	2.41	±	0.15	6.1	0.31	±	0.04	12.8
<b>Treatment 2</b>	9	3.04	±	0.09	2.9	27.2	±	0.49	1.8	2.43	±	0.04	1.7
	8	1.89	±	0.14	7.3	3.29	±	0.16	5.0	2.27	±	0.10	4.2
	7	1.35				2.87				2.30			
	6	3.21	±	0.14	4.2	7.72	±	0.30	3.9	2.05	±	0.11	5.5
	5	1.70				10.0				2.15			
	48 hr	1.71	±	0.09	5.0	10.1	±	0.10	1.0	0.37	±	0.03	7.5
<b>Treatment 3</b>	9	4.64	±	0.18	4.0	8.51	±	0.14	1.6	1.91	±	0.04	2.2
	8	3.91	±	0.12	3.2	9.69	±	0.23	2.3	1.47	±	0.04	2.6
	7	3.78	±	0.09	2.3	7.24	±	0.11	1.6	1.48	±	0.03	2.3
	6	4.09				9.18				1.35			
	5	3.32				16.2				1.41			
	48 hr	2.19	±	0.09	3.9	11.1	±	0.10	0.9	0.50	±	0.02	3.8

**Table D-5e. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Hanford Soil (Page 2 of 4)**

of added water	pH	Strontium			Yttrium			Silver		
	ppb	ppb	StD%	ppb	ppb	StD%	ppb	StD%	ppb	StD%
<b>Treatment 1</b>	9	118	± 0.81	0.7	0.20	± 0.03	16.0	<0.02		
	8	105	± 0.53	0.5	0.16	± 0.01	7.9	0.030	± 0.005	14.9
	7	94.3	± 2.26	2.4	0.53	± 0.03	6.5	<0.02		
	6	9.3	± 0.15	1.6	0.11	± 0.01	7.6	0.021	± 0.004	20.2
	5	4.6	± 0.07	1.5	0.06	± 0.02	30.8	<0.02		
	48 hr	74.5	± 1.8	2.4	0.07	± 0.01	19.2	0.036	± 0.005	14.0
<b>Treatment 2</b>	9	81.4	± 0.48	0.6		na		<0.02		
	8	92.5	± 3.5	3.8		na		0.062	± 0.006	10.0
	7	88.5				na		0.103		
	6	104	± 1.5	1.4		na		0.094	± 0.020	21.7
	5	101				na		0.061		
	48 hr	55.7	± 0.45	0.8	0.24	± 0.04	18.4	0.025	± 0.005	20.4
<b>Treatment 3</b>	9	83.4	± 1.3	1.5	6.34	± 0.17	2.7	<0.02		
	8	96.2	± 0.89	0.9	10.4	± 0.11	1.0	<0.02		
	7	101	± 0.55	0.5	5.24	± 0.18	3.3	<0.02		
	6	99.0			7.30			<0.02		
	5	110			6.65			<0.02		
	48 hr	67.8	± 0.67	1.0	0.27	± 0.03	12.0	<0.02		
		Cesium			Barium			Lanthanum		
<b>Treatment 1</b>	9	0.360	± 0.011	3.0	23.1	± 0.28	1.2	0.11	± 0.01	10.4
	8	0.549	± 0.013	2.3	21.4	± 0.43	2.0	<0.1		
	7	0.243	± 0.005	2.0	19.7	± 0.34	1.7	0.39	± 0.06	15.6
	6	0.243	± 0.001	0.5	0.85	± 0.01	1.7	<0.1		
	5	0.249	± 0.003	1.2	0.13	± 0.02	17.5	<0.1		
	48 hr	0.217	± 0.005	2.2	12.8	± 0.25	2.0	<0.1		
<b>Treatment 2</b>	9	0.217	± 0.006	3.0	14.1	± 0.21	1.5	0.19	± 0.01	3.8
	8	0.197	± 0.004	2.0	14.3	± 0.35	2.4	<0.1		
	7	0.236			13.6			<0.1		
	6	0.228	± 0.005	2.2	15.3	± 0.13	0.9	<0.1		
	5	0.230			18.9			0.13		
	48 hr	0.239	± 0.001	0.3	10.1	± 0.05	0.5	<0.1		
<b>Treatment 3</b>	9	0.235	± 0.002	1.0	13.6	± 0.13	0.9	<0.1		
	8	0.234	± 0.004	1.7	18.0	± 0.41	2.3	<0.1		
	7	0.260	± 0.003	1.2	19.1	± 0.28	1.5	0.17	± 0.06	34.2
	6	0.227			19.6			<0.1		
	5	0.234			22.1			<0.1		
	48 hr	0.240	± 0.004	1.5	14.4	± 0.12	0.8	<0.1		

**Table D-5e. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Hanford Soil (Page 3 of 4)**

	pH	Cerium			Praeseodymium			Samarium			
	of added water	ppb	ppb	StD%	ppb	ppb	StD%	ppb	ppb	StD%	
<b>Treatment 1</b>	9	0.10	$\pm$	0.02	19.0	<0.05			<0.05		
	8	<0.1				<0.05			0.07	$\pm$	
	7	0.85	$\pm$	0.03	3.7	0.10	$\pm$	0.015	0.12	$\pm$	
	6	0.15	$\pm$	0.01	7.6	<0.05			0.09	$\pm$	
	5	<0.1				<0.05			0.06	$\pm$	
	48 hr	<0.1				<0.05			0.08	$\pm$	
									0.01	17	
<b>Treatment 2</b>	9	0.25	$\pm$	0.03	12.5	<0.05			0.06	$\pm$	
	8	<0.1				<0.05			0.10	$\pm$	
	7	<0.1				<0.05			0.07		
	6	<0.1				<0.05			0.05	$\pm$	
	5	0.11				<0.05			<0.05		
	48 hr	0.13	$\pm$	0.01	11.3	<0.05			0.08	$\pm$	
									0.06	73	
<b>Treatment 3</b>	9	0.11	$\pm$	0.02	16.9	<0.05			0.10	$\pm$	
	8	<0.1				<0.05			<0.05		
	7	0.18	$\pm$	0.02	11.8	<0.05			0.07	$\pm$	
	6	<0.1				<0.05			0.05		
	5	<0.1				0.05			0.00		
	48 hr	<0.1				<0.05			0.06	$\pm$	
									0.03	55	
		Europium			Gadolinium			Dysprosium			
<b>Treatment 1</b>	9	<0.05			0.05	$\pm$	0.02	40	0.12	$\pm$	
	8	0.093	$\pm$	0.008	9	0.07	$\pm$	0.01	17	0.14	$\pm$
	7	0.069	$\pm$	0.008	11	0.13	$\pm$	0.03	27	0.15	$\pm$
	6	<0.05				0.06	$\pm$	0.01	13	0.11	$\pm$
	5	<0.05				0.07	$\pm$	0.01	20	0.17	$\pm$
	48 hr	<0.05				0.08	$\pm$	0.06	76	<0.02	
<b>Treatment 2</b>	9	0.080	$\pm$	0.017	22	0.07	$\pm$	0.03	46	0.43	$\pm$
	8	<0.05				0.06	$\pm$	0.03	51	0.09	$\pm$
	7	<0.05				0.05				0.08	
	6	0.055	$\pm$	0.019	35	0.09	$\pm$	0.02	22	0.16	$\pm$
	5	0.106				0.06				0.10	
	48 hr	<0.05				0.08	$\pm$	0.01	8	0.13	$\pm$
									0.05	40	
<b>Treatment 3</b>	9	<0.05				0.12	$\pm$	0.05	45	0.17	$\pm$
	8	0.073	$\pm$	0.023	31	0.05	$\pm$	0.04	68	0.16	$\pm$
	7	0.070	$\pm$	0.025	36	0.14	$\pm$	0.09	63	0.15	$\pm$
	6	0.076				0.11				0.18	
	5	0.100				0.08				0.10	
	48 hr	0.063	$\pm$	0.009	15	0.09	$\pm$	0.08	89	0.15	$\pm$
									0.01	9	

**Table D-5e. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Hanford Soil (Page 4 of 4)**

	pH	Holmium		Erbium			Lead		StD%
	of added water	ppb	StD%	ppb	StD%	ppb	ppb	StD%	
<b>Treatment 1</b>	9	0.024 ± 0.003	14	0.06 ± 0.02	45	<0.1			
	8	0.032 ± 0.008	26	0.04 ± 0.03	70	0.14 ± 0.032			23.7
	7	0.037 ± 0.011	31	0.07 ± 0.03	38	0.10 ± 0.023			21.9
	6	0.033 ± 0.008	25	0.09 ± 0.02	20	0.87 ± 0.014			1.6
	5	0.034 ± 0.027	78	0.03 ± 0.04	120	0.12 ± 0.002			1.3
	48 hr	<0.02		0.08 ± 0.03	42	0.13 ± 0.015			11.9
<b>Treatment 2</b>	9	0.023 ± 0.002	7	0.03 ± 0.00	9	1.04 ± 0.049			4.7
	8	0.028 ± 0.007	27	0.06 ± 0.01	14	0.11 ± 0.019			16.8
	7	0.035		<0.02		0.14			
	6	0.021 ± 0.015	71	0.04 ± 0.05	128	0.33 ± 0.022			6.7
	5	<0.02		0.03		0.17			
	48 hr	0.026 ± 0.019	73	0.04 ± 0.03	71	0.28 ± 0.005			1.7
<b>Treatment 3</b>	9	<0.02		0.05 ± 0.04	84	0.21 ± 0.014			6.5
	8	0.025 ± 0.026	103	<0.02		<0.1			
	7	0.037 ± 0.011	29	0.09 ± 0.08	90	0.14 ± 0.013			9.4
	6	<0.02		0.02		0.18			
	5	<0.02		0.02		<0.1			
	48 hr	<0.02		0.05 ± 0.02	40	0.25 ± 0.038			15.0

**Table D-5f. Inductively Coupled Plasma-Mass Spectrometry Data for the  
pH Stability Test: Hanford Groundwater (Page 1 of 4)**

of added water	pH	Chromium		Manganese		Iron		StD%
	ppb	ppb	StD%	ppb	StD%	ppb	ppb	
<b>Treatment 1</b>	9	3.1 ± 0.08	2.6	0.74 ± 0.00	0.4	217 ± 6.4	6.4	3.0
	8	0.2 ± 0.02	6.8	0.39 ± 0.02	4.6	156 ± 3.2	3.2	2.1
	7	0.3 ± 0.02	5.8	0.16 ± 0.00	2.9	90 ± 2.1	2.1	2.3
	6	0.3 ± 0.01	2.2	0.18 ± 0.01	2.9	102 ± 3.4	3.4	3.3
	5	0.4 ± 0.02	6.3	0.20 ± 0.00	1.0	100 ± 2.9	2.9	2.9
	48 h r, Sat.	0.5 ± 0.01	3.0	1.61 ± 0.02	1.1	796 ± 11.5	11.5	1.4
<b>Treatment 2</b>	9	1.9 ± 0.06	3.1	0.91 ± 0.03	3.1	243 ± 7.4	7.4	3.0
	8	2.6 ± 0.04	1.7	0.38 ± 0.00	0.2	143 ± 1.4	1.4	1.0
	7	1.1 ± 0.10	8.9	0.24 ± 0.02	7.0	104 ± 10.4	10.4	10.1
	6	0.7 ± 0.05	8.1	0.57 ± 0.01	1.3	136 ± 5.0	5.0	3.7
	5	1.2 ± 0.02	1.8	0.37 ± 0.01	3.0	122 ± 7.4	7.4	6.1
	48 hr, Sat.	1.1 ± 0.10	8.5	0.50 ± 0.03	6.2	220 ± 20.9	20.9	9.5
<b>Treatment 3</b>	9	1.3 ± 0.07	5.3	0.70 ± 0.00	0.7	169 ± 1.1	1.1	0.7
	8	0.4 ± 0.03	7.7	0.39 ± 0.00	0.7	153 ± 2.7	2.7	1.7
	7	<0.2		0.21 ± 0.01	3.1	106 ± 2.0	2.0	1.9
	6	<0.2		0.32 ± 0.00	0.8	110 ± 0.4	0.4	0.4
	5	1.8 ± 0.06	3.1	0.37 ± 0.01	3.7	149 ± 4.7	4.7	3.2
	48 h r, Sat.	<0.2		0.61 ± 0.02	3.8	331 ± 6.8	6.8	2.1
Copper								
<b>Treatment 1</b>	9	3.66 ± 0.14	3.7	5.15 ± 0.04	0.7	0.08 ± 0.01	0.01	9.0
	8	3.31 ± 0.01	0.4	5.21 ± 0.12	2.3	<0.05		
	7	0.69 ± 0.05	7.0	5.78 ± 2.28	39.5	<0.05		
	6	1.27 ± 0.09	7.0	2.64 ± 0.10	3.8	<0.05		
	5	1.44 ± 0.14	10.0	1.97 ± 0.02	1.1	<0.05		
	48 hr, Sat.	3.47 ± 0.12	3.4	7.86 ± 0.29	3.7	<0.05		
<b>Treatment 2</b>	9	2.05 ± 0.05	2.2	6.37 ± 0.20	3.1	<0.05		
	8	2.09 ± 0.39	18.8	11.4 ± 0.64	5.6	<0.05		
	7	0.76 ± 0.02	3.2	2.00 ± 0.12	6.2	<0.05		
	6	1.78 ± 0.04	2.3	5.87 ± 0.06	1.0	<0.05		
	5	1.03 ± 0.05	5.0	3.60 ± 0.26	7.1	<0.05		
	48 hr, Sat.	2.13 ± 0.17	7.9	1.23 ± 0.15	12.1	<0.05		
<b>Treatment 3</b>	9	1.13 ± 0.06	5.3	12.7 ± 0.11	0.8	<0.05		
	8	1.15 ± 0.08	7.0	2.82 ± 0.09	3.1	<0.05		
	7	0.92 ± 0.01	0.6	5.56 ± 0.09	1.7	<0.05		
	6	0.63 ± 0.03	4.8	4.80 ± 0.12	2.6	<0.05		
	5	1.32 ± 0.02	1.7	9.45 ± 0.18	1.9	<0.05		
	48 hr, Sat.	1.34 ± 0.05	3.5	11.7 ± 0.25	2.1	<0.05		

**Table D-5f. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Hanford Groundwater (Page 2 of 4)**

of added water	pH	Strontium			Yttrium			Silver		
	ppb	ppb	StD%	ppb	ppb	StD%	ppb	ppb	StD%	
<b>Treatment 1</b>	9	10.7 ± 0.16	1.5	0.22 ± 0.02	0.02	7.8	<0.02			
	8	8.2 ± 0.12	1.5	0.10 ± 0.01	0.01	12.1	0.023 ± 0.008	0.008	35.2	
	7	3.7 ± 0.05	1.4	0.09 ± 0.02	0.02	19.5	0.118 ± 0.008	0.008	6.7	
	6	4.2 ± 0.06	1.3	0.07 ± 0.02	0.02	21.9	<0.02			
	5	4.3 ± 0.12	2.8	0.23 ± 0.05	0.05	19.7	<0.02			
	48 hr, Sat.	44.6 ± 0.08	0.2	0.71 ± 0.04	0.04	4.9	0.045 ± 0.011	0.011	23.5	
<b>Treatment 2</b>	9	4.6 ± 0.08	1.7	0.21 ± 0.03	0.03	13.6	<0.02			
	8	5.6 ± 0.06	1.1	0.17 ± 0.02	0.02	14.4	0.028 ± 0.003	0.003	9.7	
	7	3.7 ± 0.16	4.4	0.12 ± 0.02	0.02	19.9	<0.02			
	6	6.2 ± 0.18	2.9	0.23 ± 0.04	0.04	15.9	0.033 ± 0.004	0.004	12.2	
	5	4.3 ± 0.02	0.4	0.28 ± 0.03	0.03	11.9	<0.02			
	48 hr, Sat.	12.2 ± 0.37	3.0	0.28 ± 0.05	0.05	17.4	0.066 ± 0.005	0.005	7.1	
<b>Treatment 3</b>	9	7.9 ± 0.04	0.5	0.31 ± 0.04	0.04	12.4	<0.02			
	8	8.7 ± 0.07	0.8	0.08 ± 0.01	0.01	12.5	0.020 ± 0.004	0.004	18.5	
	7	5.3 ± 0.08	1.6	0.10 ± 0.03	0.03	25.7	<0.02			
	6	5.3 ± 0.09	1.7	0.13 ± 0.02	0.02	18.9	<0.02			
	5	7.2 ± 0.03	0.5	0.26 ± 0.02	0.02	7.0	<0.02			
	48 hr, Sat.	17.5 ± 0.26	1.5	0.25 ± 0.01	0.01	4.4	0.035 ± 0.014	0.014	41.5	
<b>Cesium</b>										
<b>Treatment 1</b>	9	0.228 ± 0.007	2.9	1.10 ± 0.05	4.2	0.25 ± 0.05	0.05	21.5		
	8	0.236 ± 0.004	1.5	0.73 ± 0.014	2.0	<0.1				
	7	0.215 ± 0.002	1.1	0.05 ± 0.006	11.3	<0.1				
	6	0.249 ± 0.006	2.5	0.13 ± 0.012	8.9	<0.1				
	5	0.227 ± 0.005	2.1	0.13 ± 0.005	3.9	<0.1				
	48 hr, Sat.	0.228 ± 0.003	1.5	5.03 ± 0.153	3.0	0.60 ± 0.02	0.02	3.8		
<b>Treatment 2</b>	9	0.229 ± 0.002	0.8	0.23 ± 0.020	8.6	0.17 ± 0.03	0.03	15.0		
	8	0.223 ± 0.004	1.9	0.31 ± 0.019	6.2	<0.1				
	7	0.217 ± 0.006	2.9	0.08 ± 0.016	20.6	<0.1				
	6	0.211 ± 0.006	3.0	0.45 ± 0.019	4.2	0.10 ± 0.02	0.02	16.5		
	5	0.214 ± 0.009	4.4	0.18 ± 0.025	14.1	0.11 ± 0.03	0.03	29.4		
	48 hr, Sat.	0.215 ± 0.007	3.1	1.16 ± 0.024	2.1	0.17 ± 0.02	0.02	11.8		
<b>Treatment 3</b>	9	0.225 ± 0.005	2.1	0.63 ± 0.011	1.7	0.15 ± 0.01	0.01	4.4		
	8	0.213 ± 0.009	4.3	0.73 ± 0.039	5.3	<0.1				
	7	0.220 ± 0.005	2.4	0.29 ± 0.012	4.2	<0.1				
	6	0.212 ± 0.002	1.0	0.28 ± 0.011	4.0	<0.1				
	5	0.221 ± 0.003	1.5	0.55 ± 0.008	1.4	0.13 ± 0.02	0.02	13.1		
	48 hr, Sat.	0.218 ± 0.004	1.9	1.71 ± 0.040	2.3	0.16 ± 0.03	0.03	15.9		

**Table D-5f. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Hanford Groundwater (Page 3 of 4)**

of added water	pH	Cerium			Praeseodymium			Samarium			StD%
	ppb		StD%	ppb		StD%	ppb		StD%	StD%	
<b>Treatment 1</b>	9	0.49	± 0.06	13.0	0.07	± 0.026	37.4	0.10	± 0.03	27	
	8	0.15	± 0.01	8.8	<0.05			0.05	± 0.03	53	
	7	0.11	± 0.02	14.3	<0.05			<0.05			
	6	0.10	± 0.04	36.9	<0.05			0.08	± 0.04	50	
	5	0.14	± 0.01	9.2	<0.05			0.11	± 0.11	102	
	48 hr, Sat.	1.22	± 0.21	17.4	0.10	± 0.007	6.5	0.17	± 0.06	37	
<b>Treatment 2</b>	9	0.39	± 0.01	2.8	<0.05			0.07	± 0.10	135	
	8	0.14	± 0.03	20.1	<0.05			0.07	± 0.04	60	
	7	0.12	± 0.02	20.3	<0.05			0.08	± 0.08	101	
	6	0.22	± 0.03	14.3	<0.05			<0.05			
	5	0.23	± 0.03	12.0	<0.05			<0.05			
	48 hr, Sat.	0.31	± 0.08	25.2	<0.05			<0.05			
<b>Treatment 3</b>	9	0.31	± 0.04	12.3	<0.05			<0.05			
	8	<0.1			<0.05			0.08	± 0.04	46	
	7	<0.1			<0.05			0.06	± 0.08	133	
	6	<0.1			<0.05			0.05	± 0.06	122	
	5	0.16	± 0.02	10.8	<0.05			0.06	± 0.05	94	
	48 hr, Sat.	0.62	± 0.65	104.1	0.06	± 0.009	15.3	0.07	± 0.06	86	
<b>Europium</b>											
<b>Treatment 1</b>	9	0.089	± 0.012	13	0.20	± 0.07	36	0.14	± 0.03	19	
	8	<0.05			0.09	± 0.02	17	0.09	± 0.05	58	
	7	<0.05			0.18	± 0.07	41	0.04	± 0.05	142	
	6	<0.05			0.09	± 0.06	63	0.04	± 0.03	76	
	5	<0.05			0.08	± 0.03	36	0.11	± 0.06	58	
	48 hr, Sat.	0.074	± 0.009	12	1.07	± 0.09	8	0.23	± 0.05	20	
<b>Treatment 2</b>	9	<0.05			0.06	± 0.03	49	0.10	± 0.04	39	
	8	<0.05			0.06	± 0.06	106	0.07	± 0.01	19	
	7	<0.05			0.05	± 0.06	119	0.08	± 0.02	18	
	6	0.053	± 0.029	54	0.05	± 0.02	34	0.16	± 0.07	41	
	5	<0.05			0.06	± 0.03	47	0.12	± 0.05	42	
	48 hr, Sat.	<0.05			0.22	± 0.08	38	0.13	± 0.07	50	
<b>Treatment 3</b>	9	<0.05			0.12	± 0.01	12	0.14	± 0.03	25	
	8	<0.05			0.04	± 0.03	81	0.05	± 0.01	27	
	7	<0.05			0.07	± 0.04	50	0.05	± 0.01	15	
	6	<0.05			0.06	± 0.03	53	0.82	± 0.09	11	
	5	<0.05			0.08	± 0.02	23	0.21	± 0.06	30	
	48 hr, Sat.	0.055	± 0.037	66	0.84	± 0.02	3	0.08	± 0.01	16	

**Table D-5f. Inductively Coupled Plasma-Mass Spectrometry Data for the pH Stability Test: Hanford Groundwater (Page 4 of 4)**

of added water	pH	Holmium			Erbium			Lead		
	ppb		StD%	ppb		StD%	ppb		StD%	
<b>Treatment 1</b>	9	0.247	± 0.031	13	0.04	± 0.03	66	2.41	± 0.120	5.0
	8	<0.02			0.04	± 0.04	102	0.90	± 0.007	0.8
	7	0.020	± 0.005	22	<0.02			0.14	± 0.013	9.6
	6	0.022	± 0.003	15	0.03	± 0.05	174	0.11	± 0.003	2.3
	5	<0.02			0.05	± 0.01	25	0.64	± 0.034	5.2
	48 hr, Sat.	0.048	± 0.014	29	0.16	± 0.10	63	2.19	± 0.040	1.8
<b>Treatment 2</b>	9	0.071	± 0.037	51	0.07	± 0.07	98	1.01	± 0.028	2.8
	8	0.023	± 0.013	57	0.09	± 0.05	55	0.37	± 0.018	4.8
	7	<0.02			0.04	± 0.02	50	0.27	± 0.006	2.3
	6	0.021	± 0.006	28	0.03	± 0.03	90	1.21	± 0.055	4.6
	5	0.028	± 0.020	70	0.04	± 0.03	82	0.90	± 0.008	0.9
	48 hr, Sat.	0.024	± 0.006	24	0.04	± 0.06	135	0.58	± 0.017	2.9
<b>Treatment 3</b>	9	0.032	± 0.025	77	0.04	± 0.03	74	0.57	± 0.021	3.7
	8	<0.02			0.04	± 0.05	134	<0.1		
	7	<0.02			<0.02			0.75	± 0.015	1.9
	6	<0.02			0.06	± 0.01	20	0.18	± 0.015	8.2
	5	<0.02			0.04	± 0.07	189	0.85	± 0.036	4.2
	48 hr, Sat.	<0.02			0.08	± 0.02	23	0.65	± 0.014	2.1

**DISTRIBUTION**

**ONSITE (50 Copies)**

**U.S. Department of Energy  
Richland Operations Office**

---

K. M. Thompson	H0-12
A. C. Tortoso (2)	H0-12
DOE/RL Public Reading Room	H2-53

**ERC Team**

---

G. C. Henckel	H4-80
G. J. Jacksha	H9-02
M. C. Kelly	H9-03
A. J. Knepp	H4-80
T. D. LeFrancois	H9-03
T. E. Moody (30)	H9-03
S. W. Petersen	H9-03
R. P. Schroeder	H4-80
M. H. Sturges	H9-03
L. C. Swanson	H9-11
Document and Information Services (4)	H0-09
Hanford Technical Library	P8-55

**Northwest Environmental Services Testing and  
Training**

---

J. Conca	H2-52
----------	-------

BHI-00864  
Rev. 0